# Prevalence and associated factors of polypharmacy among adult Saudi medical outpatients at a tertiary care center 

Salih Bin Salih，Muhammad Yousuf，Huda Durihim，Hind Almodaimegh ${ }^{1}$ ，Hani Tamim ${ }^{2}$<br>Departments of Medicine，${ }^{1}$ Clinical Pharmacy，and ${ }^{2}$ Medical Education，College of Medicine，King Abdulaziz Medical City and King Saud Bin Abulaziz University for Health Sciences，Riyadh，Kingdom of Saudi Arabia<br>Address for correspondence：Dr．Salih Bin Salih，Department of Medicine，Division of Internal Medicine，P．O．Box 22490，King Abdulaziz Medical City，Riyadh 11426，Kingdom of Saudi Arabia．E－mail：salihbinsalih56＠gmail．com


#### Abstract

Objective：The objective of this study was to assess the prevalence of polypharmacy（ PP ）and the associated factors in medical outpatients．Materials and Methods：A cross－sectional，observational，descriptive study was carried out in adult medical outpatients attending internal medicine clinics at King Abdulaziz Medical City，Riyadh，Saudi Arabia from 1 March 2009 to 31 December 2009．PP was defined as the concomitant use of $\geq 5$ medications daily．The number of medications being currently taken by patient was recorded．Effect of patients＇age，gender，educational level，number of prescribers，disease load and disease type on PP was assessed by multivariate analysis using Statistical Package for Social Sciences Incorporated（SPSS Inc）Version 18．Results：Out of 766 patients included in the study， 683 （ $89 \%$ ）had PP． The mean number of prescribed medications，oral pills and doses was $8.8,9.6$ and 12．1，respectively．Factors significantly associated with PP included age（ $\geq 61$ years），disease load and the number of prescribers．Gender had no impact on PP while education beyond primary education significantly decreased PP．Hypertension， diabetes mellitus and dyslipidemia alone and as a cluster increased PP．Conclusion：We found an extremely high level of PP in medical outpatients at our tertiary care center．The impact of PP on medication compliance and control of underlying diseases in Saudi Arabia is unknown and needs to be studied at different levels of care．


Key words：Medical，medications，outpatients，polypharmacy

## INTRODUCTION

One of the greatest challenges facing health systems globally in the $21^{\text {st }}$ century is the increasing burden of chronic diseases．${ }^{[1]}$ The aging population with multiple chronic diseases has led to a rising prescription of medications．${ }^{[2-4]}$ As a result，the use of multiple medications or polypharmacy（PP）has become common．PP，defined as the use of multiple medications and／or the administration of more medications than is clinically indicated，often represents unnecessary use of medication．${ }^{[5]}$ However，

| Access this article online |  |
| :---: | :---: |
| Quick Response Code： |  |
| 四的F＝回 | www．jfcmonline．com |
| 2 | $\begin{aligned} & \text { DOI: } \\ & \text { 10.4103/2230-8229.121987 } \end{aligned}$ |

there is a lack of consensus regarding the actual number of concomitant medications taken by a person to be rated as PP．This is because different investigators have defined PP as the simultaneous use of two or more，${ }^{[6]}$ three or more，${ }^{[7]}$ and four or more medications．${ }^{[8]}$ Most of the recent studies from Europe and Australia have defined PP as concurrent use of five or more medications．${ }^{[9-14]}$ When guided by evidence－based－medicine，most of the medications used are rational and beneficial to the patients．Despite the differences in definition，PP is often associated with poor adherence to medications，adverse drug reactions， drug interactions，hospital admissions or readmissions， medication cascade effect and increasing costs．${ }^{[15-18]} \mathrm{PP}$ is also associated with poor nutritional status，${ }^{[10,20]}$ as well as poor clinical outcomes．${ }^{[21,2]}$

Although a previous study from a primary health care center in Riyadh，Saudi Arabia，found a PP prevalence of $21.1 \%$ as defined by the use of 4 or more medications，${ }^{[8]}$ there is little information about its prevalence and

## Salih, et al.: Polypharmacy among adult Saudi medical outpatients

associated risk factors in hospital settings. The aim of the present study was to assess the prevalence of PP and its associated factors in medical outpatients in a tertiary care setting in Riyadh, Saudi Arabia.

## MATERIALS AND METHODS

## Setting

This was an observational, cross-sectional, descriptive study of patients followed up in internal medicine outpatient clinics in the period between March 1, 2009 and December 31, 2009, at King Abdulaziz Medical City, Riyadh, Saudi Arabia. This hospital is a tertiary care teaching hospital affiliated to King Saud Bin Abdulaziz University for Health Sciences and caters for the medical needs of employees of the National Guard and their families in Saudi Arabia. After approval from the hospital research and ethical committee, patients eligible on the basis of explicit inclusion and exclusion criteria as described below participated in the study.

## Inclusion and exclusion criteria

Criteria for inclusion were: Aged 12 years or over, signed an informed consent, 2 or more visits made to the clinic and taking any prescription or non-prescription medication. The exclusion criteria included patients who were not taking any medication, diagnosed with dementia, on palliative care for any malignancy and bedridden patients on nasogastric or percutaneous endoscopic gastrostomy feeding.

## Definition of terms

PP was defined as the concurrent use of $\geq 5$ medications daily for at least 3 months. This definition was adopted from many large European and Australian studies. ${ }^{[9-14]}$ We defined prescription medications as medications prescribed by any of the doctors at this hospital or its affiliated clinics and dispensed from the affiliated pharmacies. The non-prescription medications were defined as any medication bought by the patient over the counter without a prescription. All oral, topical, inhaled and injectable medications taken on a daily basis were covered by this definition.

## Medication count

All prescription or non-prescription medications being taken by the patient were counted as a medication. These included tablets, capsules, creams, ointments, drops, syrups, liquids, suppositories, inhalers, injections and nebulized medications. Tablets and capsules were counted as pills while other medications were counted as non-pills.

For the counting of pills, a tablet or capsule was considered as one if it was used as whole or in part $(1 / 2$ or $1 / 4)$. To count the doses of medications, the medication used on
a weekly, bi-weekly or monthly basis were not included. The doses of medications taken regularly were counted as such, while the doses of Pro re nata (P.r.n) medications were taken as $50 \%$ of the maximum doses prescribed daily.

After an informed consent, eligible patients had their demographic and relevant information recorded on a special form designed for this study. This information included patients' hospital identification number, age, gender, education level, number of diseases, number of prescribers, number of drugs being used, number of doses and the number of prescription or non-prescription drugs.

Data was collected by interview, chart review and a check of all drugs being used by the patient and the tracking of their current computerized drug prescriptions. Only prescriptions filled by the patient within 1 week of being seen in the clinic were used. Patients were only included once in the study and any duplicate follow-up forms were excluded by checking the patient's unique identification number.

## Statistical analyses

The Statistical Package for Social Sciences Incorporated, Version 18 (SPSS, Inc. Chicago, Illinois, U.S.A) was used for data entry, management and analysis. Data were summarized by number and percentage or mean and standard deviation (SD), as appropriate. The association between PP and categorical variables was determined by using the Chi-square test, whereas the $t$-test was used to assess the association with continuous variables. Odds ratios (OR), $95 \%$ confidence intervals (CI) and $P$ values were calculated using a multivariate analysis. Alpha of 0.05 was used as an indication of statistical significance.

## RESULTS

During the study period, 766 patients were eligible for inclusion in the study. Of this number, 332 ( $43.3 \%$ ) were male and $434(56.7 \%)$ were female, with a mean age of 60.4 (SD 14.1) years. Most of the patients, 492 ( $64.2 \%$ ), were illiterate. The prevalence of PP was $89.1 \%$ (683 patients). Mean (SD) of the number of prescribers and number of diseases per patient was $1.69(0.97)$ and $3.86(1.39)$ respectively. The mean number, SD and Range ( R ) of prescription drugs, non-prescription drugs, number of pills, doses of pills and doses of non-pills used per patient was 8.84 (3.86, 1-24), 0.4 ( $0.275,0-5$ ), 9.56 (4.43, 1-35), 12.07 (5.90, 1-35) and 2.90 (3.12, 0-24) respectively [Table 1].

After adjustment for the confounding factors using a multivariate analysis, PP was significantly correlated with age $\geq 61$ years compared with age $\leq 60$ years (OR $6.33,95 \%$

CI 3.55-11.30, $P<0.0001$ ). Gender was not a significant factor for PP (OR 0.90, CI 0.56-1.42, $P=0.62$ ). Taking illiterate patients as a reference, PP was not significantly related with education to the primary level (OR 0.73, $95 \%$ CI 0.36-1.50, $P=0.40$ ), but it had a significant inverse relationship to secondary school level (OR 0.22, $95 \%$ CI $0.12-0.43, P<0.0001$ ) or university level of education (OR $0.12,95 \%$ CI $0.07-0.23, P<0.0001$ ) [Table 2]. PP was also significantly associated with the number of diseases, with the level of significance improving as the disease burden rose to two or more diseases. PP increased with two prescribers after which statistical significance level decreased with a further increase in prescriber numbers [Table 3]. By multivariate analysis of individual diseases, patients
suffering from dyslipidemia (DLP), hypertension (HTN), diabetes mellitus (DM), osteoarthritis, bronchial asthma, osteoporosis, heart failure and coronary artery disease had significantly higher odds of having PP than patients who did not have these diseases. Hypothyroidism and stroke were not associated with a higher PP compared with patients who did not have these diseases [Table 4]. A cluster of diseases including DLP, HTN and DM was present in $58 \%$ of the patients and was significantly associated with PP (OR 21.4, 95\% CI 8.55-53.52, $P<0.0001$ ).

## DISCUSSION

We found PP in $89 \%$ of our patients. The predictors of

Table 1: Characters of medications in patients with and without PP as mean and standard deviation

| Parameter | All patients $(\boldsymbol{n}=\mathbf{7 6 6})$ | PP $(\boldsymbol{n}=\mathbf{6 8 3})$ | No PP $(\boldsymbol{n}=\mathbf{8 3})$ | $\boldsymbol{P}$ value |
| :--- | :---: | :---: | :---: | :---: |
| Prescription drugs | $8.84(3.86)$ | $9.56(3.42)$ | $2.88(1.14)$ | $<0.0001$ |
| Non-prescription drugs | $0.04(0.28)$ | $0.04(0.28)$ | $0.05(0.27)$ | 0.79 |
| Pill burden | $9.56(4.43)$ | $10.29(4.05)$ | $3.52(2.23)$ | $<0.0001$ |
| Doses of pills | $12.07(5.90)$ | $13.00(5.47)$ | $4.47(3.24)$ | $<0.0001$ |
| Doses of non-pills | $2.90(3.12)$ | $3.17(3.15)$ | $0.65(1.51)$ | $<0.0001$ |
| PP: Polypharmcacy |  |  |  |  |

Table 2: Relationship of age, gender and educational level to PP

| Parameter | All patients ( $n=766$ ) | PP ( $n=683$ ) | No PP ( $n=83$ ) | OR | 95\% CI | $P$ value |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Age (years) |  |  |  |  |  |  |
| Mean (SD) | 60.4 (14.1) | 62.4 (12.2) | 43.7 (17.6) | - | - | <0.0001 |
| <60 (\%) | 353 (46.1) | 285 (41.7) | 68 (81.9) | 1 (Ref) | - | - |
| $\geq 61$ (\%) | 413 (53.9) | 398 (58.3) | 15 (18.1) | 6.33 | 3.55-11.30 | <0.0001 |
| Gender (\%) |  |  |  |  |  |  |
| Male | 332 (43.3) | 298 (43.6) | 34 (41.0) | 1 (Ref) | - | - |
| Female | 434 (56.7) | 385 (56.4) | 49 (59.0) | 0.9 | 0.56-1.42 | 0.64 |
| Education (\%) |  |  |  |  |  |  |
| Illiterate | 492 (64.2) | 461 (67.5) | 31 (37.3) | 1 (Ref) | - | - |
| Primary | 131 (17.1) | 120 (17.6) | 11 (13.3) | 0.73 | 0.36-1.50 | 0.4 |
| Secondary | 78 (10.2) | 60 (8.8) | 18 (21.7) | 0.22 | 0.12-0.43 | <0.0001 |
| University | 65 (8.5) | 42 (6.1) | 23 (27.7) | 0.12 | 0.07-0.23 | <0.0001 |

Table 3: Relationship of disease load and prescribers to PP

| Parameter | All patients ( $n=766$ ) | PP ( $n=683$ ) | No PP ( $n=83$ ) | OR | 95\% CI | $P$ value |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Disease load |  |  |  |  |  |  |
| Mean (SD) | 3.86 (1.39) | 4.10 (1.24) | 1.89 (0.88) | - | - | <0.0001 |
| 1 disease (\%) | 34 (4.4) | 3 (0.4) | 31 (37.3) | 1 (Ref) | - | - |
| 2 diseases (\%) | 86 (11.2) | 51 (7.5) | 35 (42.2) | 15.06 | 4.27-53.12 | <0.0001 |
| 3 diseases (\%) | 186 (24.3) | 173 (25.3) | 13 (15.7) | 137.5 | 37.02-510.82 | <0.0001 |
| $\geq 4$ diseases (\%) | 460 (60.1) | 456 (66.8) | 4 (4.8) | 1178 | 252.4-5497.7 | <0.0001 |
| No of prescribers |  |  |  |  |  |  |
| Mean (SD) | 1.69 (0.97) | 1.73 (0.99) | 1.31 (0.62) | - | - | <0.0001 |
| 1 prescriber (\%) | 423 (55.2) | 360 (52.7) | 63 (75.9) | 1 (Ref) | - | - |
| 2 prescribers (\%) | 224 (29.2) | 209 (30.6) | 15 (18.1) | 2.44 | 1.35-4.39 | 0.003 |
| 3 prescribers (\%) | 73 (9.5) | 69 (10.1) | 4 (4.8) | 3.02 | 1.06-8.57 | 0.04 |

Salih, et al.: Polypharmacy among adult Saudi medical outpatients

| Disease | Status | Patients (\%) | PP (\%) | No PP (\%) | OR | 95\% CI | $P$ value |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Dyslipidemia | No | 111 (14.5) | 65 (9.5) | 46 (55.4) | 1 (Ref) | - | - |
|  | Yes | 655 (85.5) | 618 (90.5) | 37 (44.6) | 11.82 | 7.15-19.54 | <0.0001 |
| Hypertension | No | 177 (23.1) | 124 (18.2) | 53 (63.9) | 1 (Ref) | - | - |
|  | Yes | 589 (76.9) | 559 (81.8) | 30 (36.1) | 7.96 | 4.88-12.98 | <0.0001 |
| Diabetes mellitus | No | 261 (34.1) | 195 (28.6) | 66 (79.5) | 1 (Ref) | - | - |
|  | Yes | 505 (65.9) | 488 (71.4) | 17 (20.5) | 9.72 | 5.56-16.98 | <0.0001 |
| Osteoarthritis | No | 475 (62.0) | 399 (58.4) | 76 (91.6) | 1 (Ref) | - | - |
|  | Yes | 291 (38.0) | 284 (41.6) | 7 (8.4) | 7.73 | 3.51-17.01 | <0.0001 |
| Bronchial asthma | No | 644 (84.1) | 565 (82.7) | 79 (95.2) | 1 (Ref) | - | - |
|  | Yes | 122 (15.9) | 118 (17.3) | 4 (4.8) | 4.13 | 1.48-11.48 | 0.003 |
| Hypothyroidism | No | 662 (86.4) | 586 (85.8) | 76 (91.6) | 1 (Ref) | - | - |
|  | Yes | 104 (13.6) | 97 (14.2) | 7 (8.4) | 1.78 | 0.81-4.01 | 0.15 |
| Osteoporosis | No | 686 (89.6) | 605 (88.6) | 81 (97.6) | 1 (Ref) | - | - |
|  | Yes | 80 (10.4) | 78 (11.4) | 2 (2.4) | 5.22 | 1.26-21.66 | 0.01 |
| Heart failure | No | 695 (90.7) | 614 (89.9) | 81 (97.6) | 1 (Ref) | - | - |
|  | Yes | 71 (9.3) | 69 (10.1) | 2 (2.4) | 4.55 | 1.10-18.92 | 0.02 |
| Stroke | No | 704 (91.9) | 624 (91.4) | 80 (96.4) | 1 (Ref) | - | - |
|  | Yes | 62 (8.1) | 59 (8.6) | 3 (3.6) | 2.52 | 0.77-8.23 | 0.11 |
| Coronary artery disease | No | 714 (93.2) | 632 (92.5) | 82 (98.8) | 1 (Ref) | - | - |
|  | Yes | 52 (6.8) | 51 (7.5) | 1 (1.2) | 6.62 | 0.90-48.52 | 0.03 |

PP were age, the number of prescribers, disease load and having different chronic diseases alone or together as a cluster. It was not associated with a particular gender, while a rise in the educational level beyond primary school was associated with a decrease in PP. The prevalence of PP differs in various studies depending on the definition of PP, the age of patients included and whether the patients were seen in a general practice, hospital practice or on admission. The definition of PP we used was the concomitant use $\geq 5$ medications, because this cutoff had been associated with the outcome of medication-related adverse effects for frailty, disability, mortality and falls. ${ }^{[14]}$

An $89 \%$ prevalence of PP found in our study is extremely high compared to other studies that used the same definition. For example, it was $10 \%$ in the German study of primary practice health insurance database, ${ }^{[9]}$ $33 \%$ in Denmark in a population-based general practice prescription data base, ${ }^{[10]} 46 \%$ in Italians over 65 years, ${ }^{[11]}$ $47 \%$ in Norway in admitted rheumatology and internal medicine patients, ${ }^{[12]}$ and $57 \%$ in geriatric patients of 75 years and over in Sweden. ${ }^{[13]}$ The high level of PP recorded by us may be because our study was done at a tertiary care center, where most patients are referred because of complicated diseases or multiple morbidity. It may also be because in other countries patients often have to buy drugs or pay for prescription cost per drug, whereas all medications and supplies in our center are completely free of charge. Compared to European studies, ${ }^{[11,13,15,16]}$ which showed that women used more medications than men, we did not find any significant gender difference
in the use of PP. We found the educational level of the patient to be inversely related to PP. This is in agreement with findings from Sweden, ${ }^{[13]}$ but in contrast to a Turkish study. ${ }^{[23]}$

In general, improving the educational level of the public may decrease PP by better primary prevention, healthy life-style and disease control. In agreement with other studies, ${ }^{[13,23,24]}$ we also found increasing age to be a risk factor for PP. This is not unexpected because increasing age has been linked to increased disease burden and PP. ${ }^{[25]}$ Furthermore, many of the diseases of older age are chronic in nature. PP increases with age and increasing number of chronic diseases. ${ }^{[26]}$ A cluster of diseases defined as two or more concurrent chronic diseases have been found to be associated with PP. In a study of 65 years or older patients admitted to hospitals, a cluster of diseases associated with DM was associated with PP. ${ }^{[27]}$

We also found a cluster of DLP, HTN and DM to be associated with PP. We observed a slight but statistically significant difference in PP according to the number of prescribers that often results from consultations with multiple physicians. This is in contrast to a Japanese study, ${ }^{[28]}$ that did not find any significant relationship between the number of consultations and number of medications prescribed.

Although the need to decrease PP has been stressed, longitudinal studies have shown an increasing PP over the years. ${ }^{[25,29,3]}$ Patients on PP often do not adhere to
prescribed medications. Non-adherence increases linearly with the number of medications used by a patient, being $80 \%$ with one medication compared to $20 \%$ with six or more medications. ${ }^{[3]]}$ Successful strategies to reduce PP have included home visits, ${ }^{[32,33]}$ physician feedback, ${ }^{[34]}$ physician education on the promotion of rational prescribing, ${ }^{[35]}$ medication reconciliation, ${ }^{[36]}$ and fixed dose combinations. ${ }^{[37,38]}$ The use of these strategies needs to be explored in Saudi patients.

Our study had certain limitations. The patients might not have revealed the use of over the counter medications or the medications prescribed from other hospitals. Similarly, they might not necessarily be using the medications prescribed from our hospital. The disease load might have been under-estimated since skin, ear or eye diseases were not counted. The strength of our study includes the definition and explanation of medication numbers, pills and doses prescribed to the study population as this has not been adequately addressed in the literature.

Because of the high prevalence of PP in our center, further studies in the general population, at the primary care level and in other patient groups are needed in Saudi Arabia. A study of the impact of PP on disease control, medication adherence, health-care cost, adverse drug reactions, re-admissions and falls in Saudi Arabia should also be done.

## CONCLUSION

We found a very high level of PP in the internal medicine outpatients in our tertiary care center in Riyadh, Saudi Arabia. This was related to age, educational level and number of prescribers and the burden of disease. The impact of PP on compliance to medication and control of underlying diseases in Saudi Arabia is unknown and needs to be studied at different levels of care.

## REFERENCES

1. WHO. Innovative Care for Chronic Conditions: Building Blocks for Action. Geneva: World Health Organization; 2002.
2. Anderson GF. Medicare and chronic conditions. N Engl J Med 2005;353:305-9.
3. Milton JC, Hill-Smith I, Jackson SH. Prescribing for older people. BMJ 2008;336:606-9.
4. Tinetti ME, Bogardus ST Jr, Agostini JV. Potential pitfalls of disease-specific guidelines for patients with multiple conditions. N Engl J Med 2004;351:2870-4.
5. Shoair OA, Nyandege AN, Slattum PW. Medication-related dizziness in the older adult. Otolaryngol Clin North Am 2011;44:455-71, x.
6. Veehof LJ, Stewart RE, Meyboom-de Jong B, Haaijer-Ruskamp FM. Adverse drug reactions and polypharmacy in the elderly in general practice. Eur J Clin Pharmacol 1999;55:533-6.
7. Jensen GL, Friedmann JM, Coleman CD, Smiciklas-Wright H. Screening for hospitalization and nutritional risks among
community-dwelling older persons. Am J Clin Nutr 2001;74:201-5.
8. Asiri YA, Al-Arifi MN. Polypharmacy and patterns in drug prescribing at a primary healthcare centre in the Riyadh region of Saudi Arabia. Int J Pharm Pract 2011;19:123-8.
9. Grimmsmann T, Himmel W. Polypharmacy in primary care practices: An analysis using a large health insurance database. Pharmacoepidemiol Drug Saf 2009;18:1206-13.
10. Bjerrum L, Gonzalez Lopez-Valcarcel B, Petersen G. Risk factors for potential drug interactions in general practice. Eur J Gen Pract 2008;14:23-9.
11. Nobili A, Franchi C, Pasina L, Tettamanti M, Baviera M, Monesi L, et al. Drug utilization and polypharmacy in an Italian elderly population: The EPIFARM-elderly project. Pharmacoepidemiol Drug Saf 2011;20:488-96.
12. Viktil KK, Blix HS, Moger TA, Reikvam A. Polypharmacy as commonly defined is an indicator of limited value in the assessment of drug-related problems. Br J Clin Pharmacol 2007;63:187-95.
13. Haider SI, Johnell K, Weitoft GR, Thorslund M, Fastbom J. The influence of educational level on polypharmacy and inappropriate drug use: A register-based study of more than 600,000 older people. J Am Geriatr Soc 2009;57:62-9.
14. Gnjidic D, Hilmer SN, Blyth FM, Naganathan V, Waite L, Seibel MJ, et al. Polypharmacy cutoff and outcomes: Five or more medicines were used to identify community-dwelling older men at risk of different adverse outcomes. J Clin Epidemiol 2012;65:989-95.
15. Murray MD, Kroenke K. Polypharmacy and medication adherence: Small steps on a long road. J Gen Intern Med 2001;16:137-9.
16. Colley CA, Lucas LM. Polypharmacy: The cure becomes the disease. J Gen Intern Med 1993;8:278-83.
17. Ruiz B, García M, Aguirre U, Aguirre C. Factors predicting hospital readmissions related to adverse drug reactions. Eur J Clin Pharmacol 2008;64:715-22.
18. Salazar JA, Poon I, Nair M. Clinical consequences of polypharmacy in elderly: Expect the unexpected, think the unthinkable. Expert Opin Drug Saf 2007;6:695-704.
19. Heuberger RA, Caudell K. Polypharmacy and nutritional status in older adults: A cross-sectional study. Drugs Aging 2011;28:315-23.
20. Jyrkkä J, Mursu J, Enlund H, Lönnroos E. Polypharmacy and nutritional status in elderly people. Curr Opin Clin Nutr Metab Care 2012;15:1-6.
21. Hilmer SN, Gnjidic D. The effects of polypharmacy in older adults. Clin Pharmacol Ther 2009;85:86-8.
22. Frazier SC. Health outcomes and polypharmacy in elderly individuals: An integrated literature review. J Gerontol Nurs 2005;31:4-11.
23. Gokce Kutsal Y, Barak A, Atalay A, Baydar T, Kucukoglu S, Tuncer T, et al. Polypharmacy in the elderly: A multicenter study. J Am Med Dir Assoc 2009;10:486-90.
24. Bjerrum L, Søgaard J, Hallas J, Kragstrup J. Polypharmacy: Correlations with sex, age and drug regimen. A prescription database study. Eur J Clin Pharmacol 1998;54:197-202.
25. Veehof L, Stewart R, Haaijer-Ruskamp F, Jong BM. The development of polypharmacy. A longitudinal study. Fam Pract 2000;17:261-7.
26. Slabaugh SL, Maio V, Templin M, Abouzaid S. Prevalence and risk of polypharmacy among the elderly in an outpatient setting: A retrospective cohort study in the Emilia-Romagna region, Italy. Drugs Aging 2010;27:1019-28.
27. Nobili A, Marengoni A, Tettamanti M, Salerno F, Pasina L, Franchi C, et al. Association between clusters of diseases and polypharmacy in hospitalized elderly patients: Results from the REPOSI study. Eur J Intern Med 2011;22:597-602.
28. Suzuki Y, Akishita M, Arai H, Teramoto S, Morimoto S, Toba K. Multiple consultations and polypharmacy of patients attending geriatric outpatient units of university hospitals. Geriatr Gerontol Int 2006;6:244-7.
29. Kaufman DW, Kelly JP, Rosenberg L, Anderson TE, Mitchell AA. Recent patterns of medication use in the ambulatory adult population of the United States: The Slone survey. JAMA 2002;287:337-44.
30. Linjakumpu T, Hartikainen S, Klaukka T, Veijola J, Kivelä SL, Isoaho R. Use of medications and polypharmacy are increasing among the elderly. J Clin Epidemiol 2002;55:809-17.
31. Chapman RH, Benner JS, Petrilla AA, Tierce JC, Collins SR, Battleman DS, et al. Predictors of adherence with antihypertensive and lipid-lowering therapy. Arch Intern Med 2005;165:1147-52.
32. Darnell JC, Murray MD, Martz BL, Weinberger M. Medication use by ambulatory elderly. An in-home survey. J Am Geriatr Soc 1986;34:1-4.
33. Yang JC, Tomlinson G, Naglie G. Medication lists for elderly patients: Clinic-derived versus in-home inspection and interview. J Gen Intern Med 2001;16:112-5.
34. Kroenke K, Pinholt EM. Reducing polypharmacy in the elderly. A controlled trial of physician feedback. J Am Geriatr Soc 1990;38:31-6.
35. Hogerzeil HV. Promoting rational prescribing: An international perspective. Br J Clin Pharmacol 1995;39:1-6.
36. Vawdrey DK, Chang N, Compton A, Tiase V, Hripcsak G. Impact
of electronic medication reconciliation at hospital admission on clinician workflow. AMIA Annu Symp Proc 2010;2010:822-6.
37. Bangalore S, Kamalakkannan G, Parkar S, Messerli FH. Fixed-dose combinations improve medication compliance: A meta-analysis. Am J Med 2007;120:713-9.
38. Indian Polycap Study (TIPS), Yusuf S, Pais P, Afzal R, Xavier D, Teo K, et al. Effects of a polypill (Polycap) on risk factors in middle-aged individuals without cardiovascular disease (TIPS): A phase II, double-blind, randomised trial. Lancet 2009;373:1341-51.

How to cite this article: Salih SB, Yousuf M, Durihim H, Almodaimegh H, Tamim H. Prevalence and associated factors of polypharmacy among adult Saudi medical outpatients at a tertiary care center. J Fam Community Med 2013;20:162-7.
Source of Support: Nil, Conflict of Interest: None declared.

## "QUICK RESPONSE CODE" LINK FOR FULL TEXT ARTICLES

The journal issue has a unique new feature for reaching to the journal's website without typing a single letter. Each article on its first page has a "Quick Response Code". Using any mobile or other hand-held device with camera and GPRS/other internet source, one can reach to the full text of that particular article on the journal's website. Start a QR-code reading software (see list of free applications from http://tinyurl.com/yzlh2tc) and point the camera to the QR-code printed in the journal. It will automatically take you to the HTML full text of that article. One can also use a desktop or laptop with web camera for similar functionality. See http://tinyurl.com/2bw7fn3 or http://tinyurl.com/3ysr3me for the free applications.

