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1 **BRITISH JOURNAL OF CLINICAL PHARMACOLOGY**

2 **Multi-Compartment Compliance Aids in the Community: The Prevalence of Potentially**
3 **Inappropriate Medications.**

4

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21

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23

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27 **Key Words:** Multi-compartment compliance aids, prescription drugs, drug interactions,
28 potentially inappropriate medications, social class.

29 **Abstract**

30 **Aims:** To assess the prevalence of potentially inappropriate medications (PIM) use in a
31 population of community-based multi-compartment compliance aid (MCA) users in North-
32 East Scotland.

33 **Methods:** Data for MCAs dispensed by 48 of the 50 community pharmacies in Aberdeen City
34 between 1st June to 31st October 2014, together with concurrently prescribed medications,
35 patient demographics and Carstairs Index of social deprivation were recorded. Drug-specific
36 quality indicators for PIMs from the Swedish National Board of Health and Welfare were
37 applied and bivariate logistic regression analysis used to assess for associations with
38 demographic variables.

39 **Results:** The median age was 82 years (range 12-105 years old, 59% female). A total of 1977
40 PIMs were identified affecting 57.8% of patients. A quarter of patients were prescribed ≥ 10
41 medications and 43% had a prescription containing at least one clinically significant drug-drug
42 interaction (DDI). Ten drug groups accounted for 76% of all DDIs. A significant increase in
43 the risk for at least one PIM was associated with female gender (for all indicators of PIM use),
44 age less than 80 years (three or more psychotropic medicines (OR 5.88, 2.96-11.70, $p < 0.001$)
45 and lower socioeconomic status (prescription of ≥ 10 medications (OR: 1.43, 95% CI: 1.16-
46 1.78), prescription of a long-acting benzodiazepine (OR: 1.84, CI: 1.14-2.98).

47 **Conclusions:** MCA use is associated with a significant incidence of PIMs particularly affecting
48 those under the age of 80 years and those living in deprived areas. Our findings indicate the
49 need for a more aggressive multidisciplinary approach to the review of the medications
50 prescribed to MCA users.

51

52 **What is known about this subject**

- 53 • Multi-compartment compliance aid devices are used increasingly in the UK and
54 Western Europe with the intention to maximise patient medication adherence, optimise
55 treatment benefits and minimise economic waste.

56 **What this study adds**

- 57 • Multi-compartment compliance aid use is associated with a significant number of PIMs
58 including drug-drug interactions.
- 59 • These mainly affect those under 80 years of age and those living in the most socially
60 deprived areas.
- 61 • To minimise PIM prescribing and the potential for patient harm there is a need for a
62 more aggressive multidisciplinary approach to the review of the medications prescribed
63 to multi-compartment compliance aid users.

64

65 **Introduction**

66 Multi-compartment compliance aids (MCA) are compartmentalised devices, with each discrete
67 section denoting a single dosing occasion. Formation of an MCA therefore requires
68 repackaging of solid dosage form medications, such as tablets and capsules, from the
69 manufacturer's original packaging into an MCA . The primary aim of using an MCA is to
70 maximise patient medication adherence and optimise treatment benefits. [1, 2]. However, there
71 is a lack of robust data to support the assumption that introduction of MCAs improves
72 medication adherence, as measured by pill counts and patient self-reporting [3]. Indeed, while
73 patient understanding of their own medications is widely viewed as a positive influence on
74 medication adherence [4, 5], MCA use in older people has been associated with reduced
75 knowledge of their medications, an effect that appears to be independent of patient cognitive
76 function [6].

77 Despite a lack of robust evidence, MCAs are widely employed throughout Western Europe and
78 use appears to be rapidly increasing [7-9]. Currently, there are limited data available describing
79 the prevalence of MCA use in the United Kingdom (UK).

80 While the use of MCAs is conceptually appealing to prescribers, concerns exist regarding the
81 safety of medication dispensing and the appropriateness of drug prescribing using this approach
82 [10]. The requirement to remove medications from their original packaging and insert them
83 into an MCA increases the opportunity for error within the dispensing pharmacy. Following an
84 audit of MCA dispensing in Australia, Carruthers et al reported that the medication incident
85 rate was 4.3% of issued packs with the most common causes being missing medications, supply
86 of a ceased medication, wrong strength dispensed or incorrect dosage instructions [11].

87 There is also evidence that use of MCAs is adversely associated with quality of drug
88 prescribing. Population-based studies comparing patients using an MCA with those receiving

89 routinely dispensed medications have reported that MCA use is associated with an increase in
90 PIM prescribing and potentially clinically significant drug-drug interactions (DDIs) [12, 13].
91 Belfrage et al reported recently on the results of a small study in a 100 patients using the
92 Screening Tool of Older Persons' Potentially inappropriate Prescriptions (STOPP) to assess
93 medicines issued to older patients admitted to hospital [14]. The authors reported a significantly
94 greater proportion of PIMs in patients using an MCA [14]. Similarly, in a longitudinal study of
95 older patients pre and post commencement of an MCA, Wallerstedt et al reported a sustained
96 increase in PIMs following the introduction of an MCA, which the authors postulated may be
97 related to reduced frequency of medication review once under the MCA system [15]. The
98 paucity of data supporting the use of MCAs as an aid to optimise medication adherence together
99 with data indicating increased medication incidents and poorer quality prescribing, has led to
100 growing concern over what may be seen as an increasingly untargeted approach to the use of
101 MCAs [10].

102 The majority of studies assessing PIM use in MCA users have been conducted in Scandinavia
103 and continental Europe [12, 13, 14, 15]. The aim of this study was to investigate the extent of
104 PIMs in a population of community-based MCA users in Scotland.

105 **Methods**

106 All community pharmacies (n=50) in Aberdeen City, Grampian, Scotland were sent a study
107 protocol and invitation to participate in the study by post and email with a follow-up phone call
108 from the research pharmacist one week later. Forty-eight pharmacies (96%) gave consent to
109 participate. For each MCA dispensed during the study period (1st June to 31st October 2014)
110 the following information was recorded electronically: patient demographics, medications
111 dispensed (name, strength, formulation) into an MCA, number of prescribed medications
112 dispensed out with the MCA, frequency of MCA dispensing, MCA distribution method and
113 pharmacy postal code as a surrogate for patient socioeconomic status. This information was

114 collected from patient pharmacy records, prepared MCA packs and prescriptions. Patient
115 socioeconomic status was determined using the Carstairs index score, a measure of social
116 deprivation designed originally for use in Scotland and includes factors such as employment
117 status, housing and overcrowding [16]. Patient socioeconomic status was expressed as a decile
118 of the Carstairs index score with decile 1 being the most deprived and decile 10 the least
119 deprived.

120 Because clinical data were absent and to permit international comparison, PIMs were assessed
121 using the National Indicators for Quality of Drug Therapy in Older Persons issued by the
122 Swedish National Board of Health and Welfare [13, 15, 17] as listed in Table 1. Potential DDIs
123 for medications dispensed via the MCA were assessed using the drug interaction software
124 package Lexi-Interact™ Lexicomp® [18], which classifies DDIs into 5 classes (A- no
125 interaction, B- no action needed, C- monitor therapy, D- modify regimen and X- avoid
126 combination). Only drug combinations classified as class-D or class-X interactions, both
127 denoting potential for clinically significant interaction, were recorded. PIMs and DDIs were
128 assessed by two independent researchers (Specialist Registrar in Clinical Pharmacology DC
129 and Research Pharmacist DS) and disagreements were reviewed by a third researcher
130 (Consultant Clinical Pharmacologist JSM).

131 **Statistical Analysis**

132 Binary Logistic regression analysis was used in the multivariate analysis of associations
133 between indicators of PIM and demographic variables of gender, age and Carstairs index of
134 social deprivation (expressed as odds ratio with 95% confidence intervals).

135 **Ethics Statement**

136 This study was registered as an audit with the Quality Governance and Risk Unit, NHS
137 Grampian (ID: 3044), and was therefore exempted from NHS Ethical review. Patient data was

138 anonymised at the time of data collection and stored electronically on a password-protected
139 file.

140 **Results**

141 During the study period, MCAs were issued to 2060 patients (59% female, median age 82 years
142 (IQR: 70-87), range 12 to 105 years). The majority (60.3%) of MCAs users were in the top
143 50% for socioeconomic status (Carstairs deciles 6 to 10).

144 Patients were prescribed a mean of 7.4 distinct medications per prescription (SD: 3.4, range 1-
145 23), of which, a mean of 6.4 were dispensed into an MCA (SD: 2.8., Range 1-21). Only one
146 medication was dispensed in an MCA for 2.3% (47) of the study group, while 25.1% (518)
147 were prescribed 10 or more distinct medications. Almost half of the study group (47.9%, 988)
148 had at least one medication concurrently dispensed outside of the MCA, of which 8.1% (80)
149 were prescribed five or more medications outside of the MCA. Over a fifth of the study cohort
150 (21.3%, 438) had at least a quarter of their total medications dispensed outside their MCA, and
151 4% (82) had more medications dispensed outside their MCA than within. The majority (72.1%,
152 1486) of patients had their MCA issued on a weekly basis with 0.5% (10) issued fortnightly
153 and 27.3% (563) issued monthly. Only 13.9% (n=286) of the study population collected their
154 medications in person.

155 A total of 1977 PIMs were identified in the study group, with at least one PIM occurring in
156 57.8% (1190) of the cohort, two or more in 25.1% (518) and three or more in 7.5% (n=154).

157 The maximum number of individual PIM criteria for any one patient was 5 (10 patients) and
158 the maximum total number of PIMs for a single patient was 21 caused by 12 medications (1
159 patient). The most frequent PIMs were potentially clinically significant DDIs (43.1%), 10 or
160 more distinct medications (25.1%) and medications with anticholinergic activity (16.6%). The
161 frequency of PIMs according to the individual prescribing quality indicators are reported in
162 Table 2.

163 The adjusted odds ratios for PIMs and prescribing quality indicators are reported in Table 3.
164 After adjustment for age and Carstairs index score of social deprivation, PIMs were more
165 frequently observed in females (OR 1.25, 1.04-1.51, $p < 0.05$) for all indicators of PIM, except
166 polypharmacy (10 or more medicines). PIMs of any type were more frequently observed in
167 patients under 65 years of age compared with those over 80 years (OR 1.68, 1.27-2.20,
168 $p < 0.001$). Specifically those under 65 years of age were 15 times more likely to be prescribed
169 three or more psychotropic medications (OR 15.17, 7.80-29.46, $p < 0.001$) and four times more
170 likely to be prescribed a long acting benzodiazepine (OR 4.35, 2.49-7.60, $p < 0.001$) or an
171 anticholinergic drugs (OR 3.77, 2.79-5.10, $p < 0.001$). A similar pattern was observed for those
172 aged 65-79 years with PIMs of any type being twice as likely to occur than in those over 80
173 years of age (OR 2.0, 1.6-2.53, $p < 0.001$). Specifically those 65 to 79 years of age were
174 significantly more likely to be prescribed three or more psychotropic medications (OR 5.88,
175 2.96-11.70, $p < 0.001$).

176 PIMs were significantly associated with low socioeconomic status, with those in Carstairs
177 deciles 1-5 having a 30% increased risk of a PIM of any type (OR: 1.3, CI: 1.06-1.58).
178 Specifically, polypharmacy (≥ 10 medicines) (OR: 1.43, 95% CI: 1.16-1.78), and prescription
179 for a long-acting benzodiazepine (OR: 1.84, CI: 1.14-2.98).

180 A total of 1359 potentially clinically significant DDIs were identified with 43.1% (887) MCA
181 users having at least one DDI. Medications from 33 different drug groups were involved in
182 potentially clinically significant DDIs. The maximum number of potentially clinically
183 significant DDIs recorded for a single patient was 19 caused by 12 medications. DDIs were
184 more likely to occur in those with polypharmacy (>10 prescription medications in MCA) (3.95,
185 3.18-4.92, $p < 0.001$), females (1.29, 1.07-1.55, $p < 0.01$) and those aged 65 to 79 years olds (1.62,
186 1.31-2.02, $p < 0.001$). The ten top drug groups accounting for 72.7% of DDIs were
187 antidepressants (13.9%), calcium supplements (9.2%), statins (8.5%), antiplatelets (7.9%),

188 proton pump inhibitors (6.9%), anticonvulsants (6.1%), antihypertensive agents (6.0%).
189 antipsychotics (5.6%), levothyroxine (5.0%) and neuropathic analgesics (3.6%).

190 **Discussion**

191 This is the first study in the UK to report the prevalence of PIMs in a population of MCA users
192 in the community. Over half of the patients issued with an MCA had at least one PIM and more
193 than two fifths at least one potential clinically significant DDI. While previous studies have
194 reported similar levels of PIM, the rate for potentially clinically significant DDIs observed in
195 our study are five-fold greater than the 8-9% reported for an older Swedish population [12, 13].
196 The reasons for the apparent increase in prevalence of DDIs is unclear but may be due to the
197 wider use of medications such as psychotropic medications that are particularly associated with
198 DDIs in the relatively younger population seen in this study [12, 13].

199 The adjusted odds ratio for all the indicators for PIMs were increased in those under the age of
200 65 years compared to those aged ≥ 80 years, particularly for use of \geq three psychotropic
201 medications and long-acting benzodiazepines, possibly reflecting the nature of the disease
202 burden (mental health issues) in the under 65 year age group necessitating MCA use. Of interest
203 is the observed increase in the adjusted odds ratio for all but one of the indicators for PIMs in
204 those aged 65-79 years relative to those ≥ 80 years. This observation that has been previously
205 reported by others and is believed to be due to the healthy survivor effect in those ≥ 80 years
206 of age [12, 19]. Nonetheless, these findings indicate the need to focus particular attention on
207 prescribing in MCA users under the age of 80 years.

208 To the best of the author's knowledge socioeconomic status has not been included in previous
209 studies reporting medication safety in MCA users. A significant relationship was observed
210 between social deprivation and PIM occurrence in the lowest socioeconomic groups, in
211 particular polypharmacy or a prescription for a long-acting benzodiazepine. It is well
212 recognised that individuals of lower socioeconomic status tend to experience worse health and

213 higher levels of anxiety and it is possible that these observations reflect an increased disease
214 burden [20, 21].

215 Unavoidably, a proportion of MCA users (almost half of our study population) require
216 medications such as inhalers, which are not compatible with dispensing into an MCA.
217 However, our finding that over a fifth of the study population had more than a quarter and
218 almost one in twenty had more than half of their medications dispensed outwith an MCA
219 detracts from the simplicity of application and the goal of improved adherence, which MCAs
220 are intended to achieve [15].

221 There is an increased prevalence of both cognitive impairment and renal dysfunction amongst
222 MCA users, indicating a higher burden of disease in this patient population [14]. It is therefore
223 unsurprising that only 14% of the patients in this study collected their prescriptions in person.
224 However, missing this opportunity for direct pharmacist-patient interaction may be significant
225 since regular interaction between pharmacists and patients has been associated with improved
226 medication adherence [22]. Our finding that more than two fifths of subjects were exposed to
227 a potential DDI further reinforces the importance for the pharmacist and prescribing physician
228 to collaboratively assess both the MCA user and their prescription on a regular basis.

229 There is little data regarding the prevalence of MCA use in the UK, however in 2001, Nunney
230 et al estimated that there were 100,000 MCA users in the UK, equating to a 170/100,000 of the
231 population [23]. Our data suggest that the prevalence of MCA use in 2015 is now 900/100,000
232 of the population, representing a greater than five-fold increase over a 14 year period, which
233 appears disproportionate to the 1.2 fold increase in the UK older population over the same
234 period [24, 25].

235 **Study Strengths and Weaknesses**

236 Although this study provides insight into medication use by MCA users under 65 years of age,
237 the criteria used were originally validated in an older population (>65 years) and therefore may

238 not be fully generalisable to all age groups [17]. However, it may be argued that the PIM criteria
239 are equally applicable to all age groups and the presence of morbidity and comorbidity may be
240 more relevant than age *per se*.

241 Our finding that socioeconomic status appears to be independently associated with PIMs is
242 significant, however we did not directly account for patient disease burden which is also
243 directly associated with socioeconomic status [20]. Therefore, the observed relationship
244 between socioeconomic status and PIM may be largely accounted for by disease burden.
245 Patient socioeconomic status was determined from the supplying pharmacy postcode, thus
246 assuming that both patient and pharmacy lay within the same geographical area. It has been
247 reported that almost 90% of patients live within 1.6 kilometres of their pharmacy suggesting
248 that this is a reasonable assumption to make [26]. The study population were exclusively
249 residents of the North East of Scotland and hence its findings may not be generalisable to the
250 whole UK population and beyond.

251 The lack of clinical data prevented the use of more comprehensive screening tools for
252 inappropriate medicine use such as the STOPP and START criteria, which prevented
253 assessment of potential prescribing omissions and clinically relevant inappropriate medicine
254 use. Therefore, our results are likely to be an underestimation of the actual PIM prevalence.

255 **Conclusions**

256 A significant proportion of MCA users in this study were prescribed PIMs including DDIs,
257 with those under the age of 80 years and those living in the poorest areas at greater risk. The
258 simplification of medication consumption, which the MCA is designed to provide, appears to
259 be confounded in a significant number of individuals by the concurrent supply of medications
260 outwith the MCA system. Our findings indicate a need for a more aggressive multidisciplinary
261 approach (involving prescriber, dispensing pharmacist and patient) to the review of the

262 medications prescribed to MCA users, which is particularly poignant given the apparent
263 increase in MCA use in the UK.

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273 None

274 **Conflict of Interest/Disclosure:** The authors have no conflicts of interest, financial or
275 otherwise, to declare.

276 **Author Contributions:**

277 **DC:** Designed the study, collected data, analysed data and wrote the manuscript.

278 **JM:** Collected data, analysed data and wrote the manuscript.

279 **DS:** Designed the study, analysed data and wrote the manuscript.

280 **JSM: Principal Investigator.** Designed the study, analysed data and wrote the manuscript
281 and acts as guarantor for the study.

282

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357

358 **Legend for Tables**

359 **Table 1:** Indicators of Potentially Inappropriate Medicines with Qualifying Drug Classes.

360 Presence of a PIM was dependent solely on the prescription of a qualifying medication
361 regardless of preparation, dose or indication. (ATC Denotes Anatomical Therapeutic Chemical
362 WHO Classification System).

363

364 **Table 2:** Prevalence of Potentially Inappropriate Medicines Associated with MCA (n=2060)

365

366 **Table 3:** Adjusted Odds Ratios For Potentially Inappropriate Medicine Use According To
367 Prescribing Quality Indicators, Adjusted for Age, Gender, Residence And Carstairs Index
368 Score.

369 (NS denotes variable-indicator combinations that were not significant in the multivariate analysis model. *
370 denotes p<0.05 relative to reference group within variable category. ** denotes p<0.01 and *** denotes
371 p<0.001. LA Benzo = long-acting benzodiazepine. Any PIM = presence of at least one indicator for potentially
372 inappropriate medicine, ref = reference variable.)