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9 **Prevalence and potential interactions from concurrent use**
10 **of prescription drugs, herbal medicinal products and**
11 **dietary supplements among older adults: a cross sectional**
12 **survey**
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31 **Abstract**

32 **Background**

33 Polypharmacy is common among older adults, with increasing numbers also using
34 prescription drugs with herbal medicinal products (HMPs) and dietary supplements.
35 There is no reliable evidence from the UK on concurrent use of HMPs and dietary
36 supplements with prescription drugs in older adults.

37 **Aim**

38 To establish prevalence of concurrent prescription drugs, HMPs and dietary
39 supplements among UK community dwelling older adults and identify potential
40 interactions.

41 **Design and Setting**

42 Cross sectional survey of older adults registered at two general practices in South
43 East England.

44 **Methods**

45 A questionnaire asking about prescription medications, HMPs and
46 sociodemographic information posted to four hundred older adults ≥ 65 years,
47 identified as taking one or more prescription drug.

48 **Results**

49 Respondents = 155, response rate = 39%, prevalence of concurrent HMP and
50 dietary supplements with prescriptions = 33%. Females more likely than males to be
51 concurrent users (43.4% versus 22.5%; $p = 0.009$). Number of HMPs and dietary
52 supplements ranged from 1 to 8, (mean = 3, median = 1; SD = 1.65). The majority of
53 concurrent users (79.6%) used dietary supplements with prescription drugs. Most
54 commonly used dietary supplements were cod liver oil, glucosamine, multivitamins
55 and Vitamin D. Others (31.6%) used only HMPs with prescription drugs. Common

56 HMPs were evening primrose oil, valerian, Nytol herbal[®] (a combination of hops,
57 gentian and passion flower). Sixteen participants (32.6%) were at risk of potential
58 adverse drug interactions.

59 **Conclusion**

60 General Practitioners should routinely ask questions regarding herbal and
61 supplement use, to identify and manage older adults at potential risk of adverse-drug
62 interactions.

63

64 **How this fits in**

- 65 • Older adults use herbal medicines and dietary supplements with prescription
66 drugs to manage chronic conditions and to maintain health.
- 67 • The use of HMPs and dietary supplements with prescription drugs among
68 older adults is under researched.
- 69 • Concurrent use of HMPs and dietary supplements with warfarin, statins and
70 anti-inflammatory drugs is common among UK older adults.
- 71 • One in three concurrent user is at risk of a potential herb-drug or supplement-
72 drug interaction.

73

74 **Keywords**

75 general practice; herbal medicine; dietary supplements; herb-drug interactions;
76 polypharmacy

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80 **INTRODUCTION**

81 Polypharmacy is a recognised patient safety risk among older adults.^{1 2} Although
82 standard definitions of polypharmacy do not generally include herbal medicinal
83 products (HMPs) and dietary supplements, they increase the risk of adverse drug
84 events through potential interactions.³⁻⁵ Use of herbal medicines and dietary
85 supplements by older adults is common.^{6-8 8-10} Also, medication-related problems are
86 higher among older adults because of co-morbidities and reduced clearance of
87 pharmacologically active compounds.¹¹

88

89 A recent systematic review¹² established that concurrent use of prescription drugs
90 and HMPs is substantial among older adults, with potential interactions from some
91 herb-drug combinations such as garlic/aspirin and ginseng/warfarin. However, the
92 only available UK study on this issue among older adults¹³ is nearly 15 years old,
93 included people 50 and over and relied on a self-selected sample from a magazine's
94 readership.

95

96 The aims of the study were to establish prevalence, identify patterns and potential
97 herb-drug interactions from concurrent use of prescription drugs, HMPs and dietary
98 supplements among UK community dwelling older adults.

99 **METHODS**

100 We conducted a cross-sectional survey using self-administered questionnaires
101 between January and April 2016. Data was from a purposive sample of community
102 dwelling older adults registered at two general practices in South East England.
103 Practice A; small village practice in Essex, with predominantly White population and

104 about 19.5% aged ≥ 65 years. Practice B; large urban practice in North London with
105 sizeable British, Black, Asian and minority ethnic (BAME) population, up to 9% aged
106 ≥ 65 years.

107

108 Ethics review and support were obtained from the London-Hampstead Research
109 Ethics Committee (15/LO/1870).

110

111 Informed by findings from a systematic review on concurrent use of HMPs with
112 prescription drugs,¹² the questionnaire covered demographics, prescription drugs,
113 HMPs and dietary supplements not prescribed by the GP, the rationale for taking
114 them and how often. Participants were also asked about side effects and perceived
115 benefits of medicines and/or HMPs (Appendix A).

116

117 As no consistent terms exist for HMPs, some examples of common HMPs were
118 included to show participants the possible breadth of responses. Questionnaire was
119 piloted among 15 community dwelling older adults. Following minor amendments,
120 the questionnaire consisted of 18 questions and took between 20 to 30 minutes to
121 complete.

122

123 **Sample size calculation and sampling**

124 This was based on previous research evidence that up to 50% of older adults use
125 HMPs.¹⁴⁻¹⁶ To achieve a confidence interval (CI) of 95% and $\pm 7.5\%$ accuracy, a
126 power calculation (using G*Power version 3.1.9.2) gave total sample size required
127 as 171 respondents (using the worst -case scenario of actual proportion being 50%).

128

129 Purposive sampling^{17 18} was used to obtain a sample of older adults registered at
130 two general practices in Essex and Haringey. Participants had to be ≥65 years, using
131 at least one prescription drug and able to consent. Terminally ill patients, patients
132 with dementia or assessed to lack capacity were excluded.

133

134 To account for attrition from low responses which are common with mailed surveys,
135 two hundred participants who met inclusion criteria were randomly selected from a
136 list of patients in each of the two practises. Practice administrators posted study
137 packs (letter of invitation addressed personally to the patient, participant information
138 sheets (PIS), questionnaire and a reply-paid envelope) to selected patients.

139 Reminder letters were sent after two weeks. If there was no reply, new study packs
140 were sent to all non-responders.

141

142 **Data analysis**

143 Respondents were defined as 'concurrent users' if they reported using HMPs or
144 dietary supplements. Anonymised data were recorded, double checked;
145 discrepancies corrected and analysed using SPSS version 23.0. Descriptive
146 statistics were calculated to summarise the sample. Using Fisher's exact test,
147 associations between concurrent use and demographic factors i.e. sex, educational
148 qualifications and living arrangements was tested. Statistical significance was
149 measured at 5% level.

150

151 Potential interactions between combinations of prescription drugs, HMPs, and
152 dietary supplements were assessed using *Stockley's Herbal Medicines Interactions*

153 <https://www.medicinescomplete.com/mc/shmi/current/>. *Stockley's Herbal Medicines*
154 *Interactions* is a comprehensive evidence database. Information regarding potential
155 interactions between herbal medicines (including nutritional supplements and some
156 food) and conventional medicines are expertly assessed with practical advice
157 provided and regularly updated.

158

159 Interactions are rated on three criteria; whether action is required to address the
160 interaction (action), likely effect of the interaction on the patient if unmanaged
161 (severity) and weight of available evidence regarding the interaction (evidence).

162 Based on these criteria, drug-herb or drug-supplement combination is described by
163 one of five outcomes ranging from no interaction, doubt about outcomes, potentially
164 hazardous outcomes, significant hazard and life threatening outcomes.

165

166 All drug-herb and drug-supplement combinations were assessed and rated for
167 potential interactions by TA and double checked by SK (a registered pharmacist).

168 Any disagreements on rating were resolved through discussions.

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176 **RESULTS**

177 Table 1 shows sociodemographic characteristics of the 155 respondents, a response
178 rate of 39%. Power calculation based on 80% power, and significance level of 5% for
179 a two-tailed test showed a sample of 128 (64 in each group) was sufficient to detect
180 medium standardised effect size and thus the results are unaffected by the shortfall
181 in the desired number of respondents.

182

183 Of these 155 respondents, 81 (53%) were females, 72 (47%) males; two participants
184 did not indicate gender. The majority (109; 70.3%) were 65-74years, none was ≥ 95
185 years. About half of participants (75; 48.4%) lived with a partner or spouse while
186 31.6% lived alone. Most respondents identified as White (132; 86.3%), with 15
187 (9.6%) Black or Black British and only one Asian or Asian British.

188

189 There was almost an even split in educational level between respondents; 71
190 (45.8%) with further education after secondary school and 77 (49.6%) with no
191 education beyond secondary school. Only one participant had a clinical background
192 in nursing.

193 **Prevalence and pattern of concurrent use**

194 Six responders provided no information on medications; therefore 149 were included
195 in the final analysis yielding a useable response rate of 37%. Of these, 49 or
196 approximately one in three reported concurrent use of HMPs and dietary
197 supplements with prescribed medications in the last 12 months. The prevalence of
198 concurrent use is thus 33%.

199

200 Females were more likely than males to be concurrent users (43.4% versus 22.5%,
201 $p = 0.009$). There is little difference between groups that had or had no further
202 education with each having about a third reporting concurrent use. Similarly, there
203 were little differences between the categories of living arrangements. Those living
204 with children reported somewhat lower levels of concurrent use but this was not
205 statistically significant.

206 **Table 1 here**

207

208 Concurrent users ($n=49$) reported 55 herb-drug and supplement-drug combinations.
209 The number of HMPs and dietary supplements ranged from 1 to 8, mean value of 3
210 (standard deviation = 1.65, median = 1).

211

212 All respondents were taking one or more prescription drugs on a regular basis, range
213 1 to 18 (median = 3). A total of 180 different types of prescription drugs reported by
214 149 participants (mean = 3.96, SD = 2.52). The most reported drug classes were
215 statins (69; 46.3%), beta blockers (26; 17.4%), calcium channel blockers (23;
216 15.4%), non-steroidal anti-inflammatory drugs (NSAIDs) (19; 12.7%), biguanides (18;
217 12.1%), angiotensin-converting enzyme (ACE) inhibitors (18; 12.1%) and proton
218 pump inhibitors (18; 12.1%).

219

220 Ten concurrent users (31.6%) used only HMPs with prescriptions. Thirty six different
221 herbs (used either singly or as a combination product) were reported. The most
222 commonly used HMPs were evening primrose oil, valerian, Nytol herbal[®]
223 (combination of hops, valerian, gentian and passion flower), garlic, cinnamon and
224 Echinacea (Figure 1).

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Figure 1 here

228 The majority of concurrent users (39; 79.6%) were taking dietary supplements,
229 including vitamins and minerals. The most combined dietary supplements were cod
230 liver oil, glucosamine, multivitamins and Vitamin D (Figure 2). Of the 49 concurrent
231 users; 13 (26.5%) reported using both HMPs and dietary supplements concurrently
232 with prescription drugs. 38.8% of concurrent users used three or more HMPs or
233 dietary supplements concurrently with prescription drugs.

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Figure 2 here

241 **Potential interactions between HMPs, dietary supplements and**
242 **prescription drugs**
243

244 Just over half of the 55 herb-drug and supplement–drug combinations (n= 28,
245 50.9%), were assessed as ‘no interaction’ or ‘no interaction of clinical significance’.
246 However, 21 combinations were categorised as interactions with ‘doubts about the
247 outcome of concurrent use’. Three combinations were rated as ‘potentially
248 hazardous’ and three of ‘significant hazard’ (Table 3).

249

250 The HMPs implicated in the potential risk for interaction include:

- 251 • flaxseed
- 252 • evening primrose oil
- 253 • St John’s wort
- 254 • peppermint,

- 255 • Senna
- 256 • Echinacea
- 257 • Hawthorne
- 258 • Nytol herbal[®] (combination of Hops, Valerian, Gentian, passion flower)
- 259 • Ginkgo.

260

261 The five dietary supplements implicated are:

- 262 • glucosamine
- 263 • cod liver oil
- 264 • omega 3 fish oil
- 265 • calcium carbonate
- 266 • a multivitamin.

267

268 A majority of the identified interactions involved potential alterations in the
269 concentration or effect of the prescription drugs, including calcium channel blockers,
270 HMG-CoA reductase inhibitors (statins) and aspirin. The seven herb-drug and
271 supplement–drug interactions assessed as having the potential for hazardous
272 outcomes, relates to increase in blood-glucose concentrations, risk of bleeding and
273 reduced efficacy or bioavailability of the prescription drug.

274

275

276 **Table 2 here**

277

278 **DISCUSSION**

279 **Summary**

280 Almost one-third of older adults (33%) in our sample were using a HMP or
281 supplement concurrently with prescription drugs. About one in three concurrent
282 users was at risk of a potential herb-drug or supplement-drug interaction. If applied
283 to the UK population that would mean 3.85 million older adults in the UK are at risk
284 of at least one potential herb-drug or supplement-drug interaction. Importantly, 6
285 combinations (Table 2) have potential for hazardous outcome or significant hazard.

286

287 **Strengths and limitations of the study**

288

289 To our knowledge, this is the first UK study in over 10 years assessing concurrent
290 use of prescriptions, HMPs and dietary supplement among older adults.¹³ Our study
291 participants were community dwelling older adults recruited from general practices,
292 previous UK studies have researched patients with cancer,¹⁹ diabetes^{20 21} and
293 pregnant women.²²

294

295 A response rate of 39% is disappointing but high for a study of this kind and enough
296 to provide credible findings. Examples of HMPs and dietary supplements were
297 included in the questionnaire, it is possible that some participants did not consider
298 products such as garlic or ginger used for medicinal purposes as HMPs and did not
299 report them.

300

301 **Comparison with existing literature**

302 The prevalence of concurrent prescription, HMPs and supplement use among older
303 adults varies from country to country, between 9% and 88.3%.²³⁻³³ The prevalence
304 of 33% reported in this study is similar to studies in North America.²³ However, lower
305 prevalence of 15%²⁷ and 22.8%²⁸ were reported in two other US studies. Variation
306 in prevalence could be explained by different definitions and inclusion criteria for
307 HMPs and level of detail required about HMPs. Not all studies focused exclusively
308 on older people and some were more successful; recruiting people from BAME
309 backgrounds.

310

311 Another possible explanation for discrepancies in prevalence of concurrent use in
312 our study and those in the literature may be problem of recall. Some studies asked
313 participants to recall products using self-completed questionnaires.^{13 28-30 34} Others
314 interviewed face-to-face, checked and recorded medications.^{26 27 31 35 36} Some
315 patients do not consider HMPs and dietary supplements as medicines. Even when
316 questioned, they may not always remember to disclose them. This highlights the
317 need for direct questioning and for healthcare practitioners to ask about use of those
318 HMPs and supplements by name, which this research has shown to be most at risk
319 of interactions.

320

321 All older adults in our sample were using at least one prescription drug, the range of
322 prescription drugs was similar to those reported by older adults in a previous US
323 study.³⁷ The number of prescription drugs taken ranged from 1 to 18, with 35% of our
324 sample taking 5 or more prescription medications. This is consistent with findings of
325 a study among US adults³⁸ and UK data.³⁹ The number of HMPs and supplements

326 used by our participants ranged from 1 to 8, and unexpectedly almost half of the
327 concurrent users were using three or more HMPs and supplements.

328

329 Certain demographic and clinical characteristics are associated with concurrent
330 medicine use. Women,^{13 36} older age groups,³⁵ people with chronic conditions such
331 as diabetes and high blood pressure,^{3 31} people with less than high school education
332 ^{31 40} and people on low income ⁴⁰ are more likely to be concurrent users. Our finding
333 that 43.4% females were concurrent users compared with 22.5% males, confirms
334 previous findings that women tend to use more herbal and dietary supplements.^{13 14}
335 ³⁶ The increased odds for a co-user to be female (34% vs 18%, $p=0.001$) has been
336 previously reported.³⁴ This is likely to be for many reasons; women generally tend to
337 live longer than men,^{41 42} they are the main carers for children and older people,⁴³
338 tend to buy medicines and remedies for the home, and also tend to use more weight
339 loss products than men.⁴⁴

340

341 Age was not significantly associated with concurrent use in our study, and in two
342 other similar studies.^{31 32} However, concurrent use was highest among our
343 participants in the age group 65-74 years (34.9%) but declined among those 85
344 years and older. Arcury et al ⁴⁵ reported significant association between age and use
345 of herbal remedies but observed a similar trend of decreased use among those ≥ 75
346 years and older.

347

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349

350 **Implications for research and practice**

351 This research has highlighted potential risk of interactions with certain combinations
352 of prescription drugs, HMPs and dietary supplements. Therefore, healthcare
353 professionals should routinely ask questions regarding use of other medications that
354 are not prescribed. The problem with recall and what patients think are
355 HMPs/supplements or not needs to be addressed. There are stages in the process
356 of prescribing and dispensing which could also be optimised, such as printing a
357 warning on prescriptions or looking at counselling given by pharmacists on
358 dispensing. Targeted questioning about use of any alternative medicine or
359 supplements could initiate conversations about wider HMP use and possible
360 interactions. Liaising with community pharmacists could also raise awareness of a
361 potential problem, particularly for older people on warfarin and statins.

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379

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384

385 **Conflicts of Interest**

386 The authors have declared no conflicts of interest directly relevant to the content of
387 this study.

388

389 **Authors' Contributions**

390 This study is part of Taofikat Agbabiaka's PhD in Health Research at the University
391 of Hertfordshire. All authors were involved in the study design. TA undertook the
392 data management and primary analysis, and wrote a first draft of the manuscript. NS
393 supervised data analyses. SK reviewed and validated potential interactions with TA.

394 All authors contributed to further drafts and approved the final manuscript.

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Table 1: Demographics of concurrent users and non-concurrent users (n=149)

Demographics		Concurrent users n = 49 (%)	Non-Concurrent users n = 100 (%)	Total	P value (Fisher's Exact test)
Age group	65-74	37 (34.9)	69 (65.1)	106 (100)	p = 0.729
	75-84	11 (33.3)	22 (66.7)	33 (100)	
	85-94	2 (20.0)	8 (80.0)	10 (100)	
Sex	Male	16 (22.5)	55 (77.5)	71 (100)	p = 0.009
	Female	33 (43.4)	43 (56.6)	76 (100)	
Ethnic background	White	40 (31.7)	86 (68.3)	126 (100)	p = 0.184
	Asian or Asian British	1 (100.0)	0 (0.0)	1 (100)	
	Black or Black British	5 (33.3)	10 (66.7)	15 (100)	
	Mixed/Multiple background	1 (33.3)	2 (66.7)	3 (100)	
	Other Ethnic Group	2 (100.0)	0 (0.0)	2 (100)	
Further education	Yes	24 (34.3)	46 (65.7)	70 (100)	p = 0.862
	No	25 (32.5)	52 (67.5)	77 (100)	
Living arrangement	Alone	17 (37.0)	29 (63.0)	46 (100)	p = 0.929
	With partner/spouse	26 (35.6)	47 (64.4)	73 (100)	
	With partner/spouse and children	4 (26.7)	11 (73.3)	15 (100)	
	With children	2 (28.6)	5 (71.4)	7 (100)	
	Other	1 (20.0)	4 (80.0)	5 (100)	

Bold values shows statistically significant result (p < 0.05)

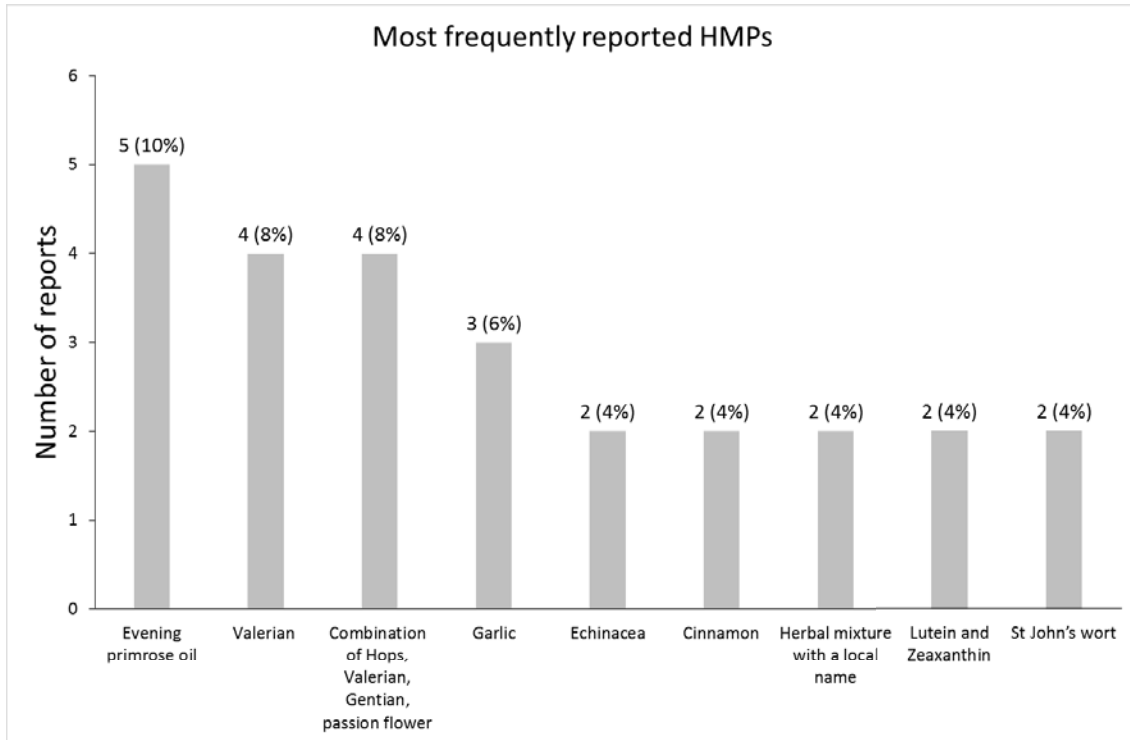


Figure 1 HMPs most frequently used concurrently with prescription by UK older adults

*Percentages sum to more than 100% as individuals could report more than one HMP.

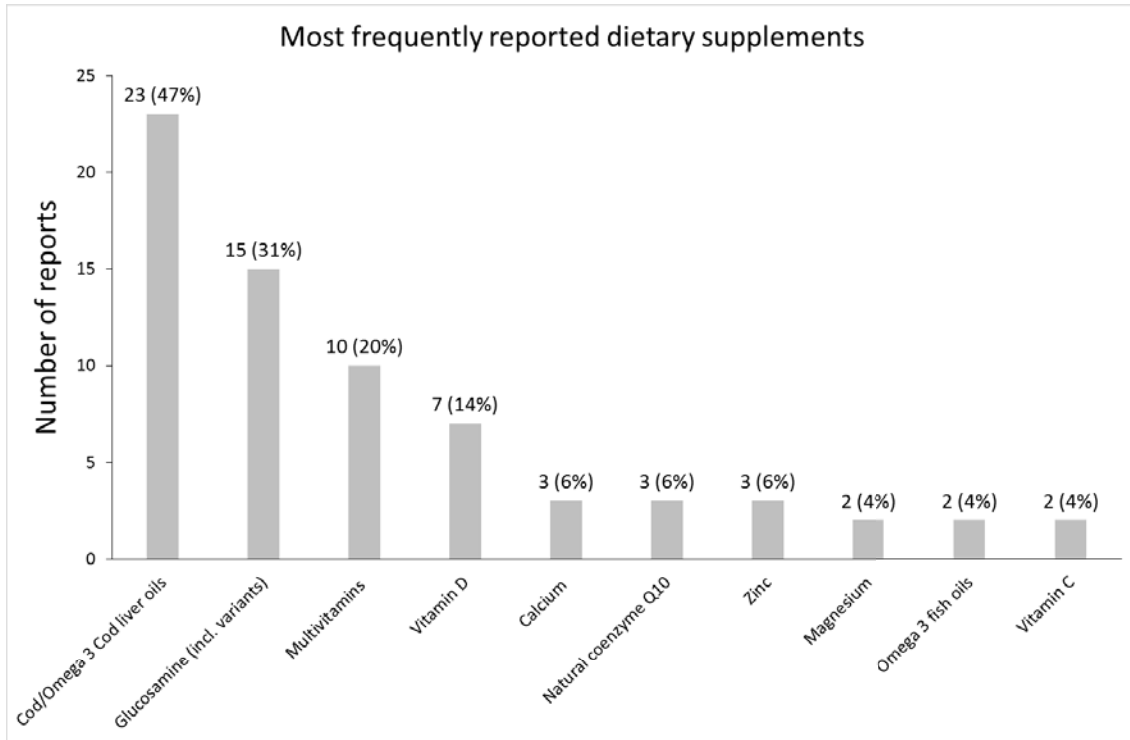


Figure 2 Supplements that are most frequently used concurrently with prescription by UK older adults

*Percentages sum to more than 100% as individuals could report more than one HMP.

Table 2 Evaluation of potential interactions from HMPs, dietary supplements and prescription drugs

HMPs/ Dietary Supplement	Prescription medicine [no of patients ^a]	Possible interactions *
HDI Interaction category : significant hazard, dosage adjustment or close monitoring is needed		
Bonecal	Levothyroxine	The efficacy of levothyroxine has been reduced by calcium carbonate. Calcium acetate and calcium citrate reduced levothyroxine absorption in pharmacokinetic studies
Peppermint	Lansoprazole	Antacids may compromise the enteric coating of some commercially available peppermint oil capsules. H ₂ -receptor antagonists and proton pump inhibitors may interact similarly.
St John's wort	Amlodipine	St John's wort significantly reduces the bioavailability of verapamil. Other calcium-channel blockers would be expected to interact similarly.
HDI Interaction category : A potentially hazardous combination		
Glucosamine	Metformin	In a controlled study, glucosamine supplements with chondroitin had no effect on glycaemic control in patients taking oral antidiabetic drugs, but increases in blood-glucose concentrations have occurred in patients with treated and untreated diabetes.
Omega 3 fish oil	Bisoprolol [2]	The hypotensive effect of propranolol might be enhanced by fish oils.
Ginkgo	Rabeprazole	Ginkgo Modestly reduces omeprazole levels. Most other proton pump inhibitors are likely to be similarly affected.
HDI Interaction category : Doubt about outcome of concurrent use ^b		
Omega 3 fish oil	Aspirin [2]	The concurrent use of aspirin and fish oils caused at least additive effects

		on bleeding time in healthy subjects, but clinical studies in patients taking aspirin alone and with clopidogrel have found no evidence of an increase in incidence of bleeding episodes.
Cod liver oil	Aspirin [2]	The concurrent use of aspirin and fish oils caused at least additive effects on bleeding time in healthy subjects, but clinical studies in patients taking aspirin alone and with clopidogrel have found no evidence of an increase in incidence of bleeding episodes.
Cod liver oil	Bisoprolol Propranolol	The hypotensive effect of propranolol might be enhanced by fish oils.
Flaxseed	Rivaroxaban	Limited evidence suggests that flaxseed oil may have some antiplatelet effects, which could be additive with those of conventional antiplatelet drugs, and increase the risk of bleeding with anticoagulants.
Green tea	Lisinopril	Both black and green tea might cause a modest increase in blood pressure, which might be detrimental to the treatment of hypertension. Green tea reduced the effects of nadolol on blood pressure in healthy subjects.
Senna pods	Indapamide	Theoretically, patients taking potassium-depleting diuretics could experience excessive potassium loss if they also regularly use, or abuse, anthraquinone-containing substances such as <i>Senna</i> .
Glucosamine	Co-codamol Paracetamol	Limited evidence suggests that glucosamine might reduce the efficacy of paracetamol (acetaminophen).
Glucosamine	Furosemide Bendroflumethiazide [2]	Limited evidence from a large open study suggests that unnamed diuretics might slightly reduce the efficacy of glucosamine to some extent.
Echinacea	Simvastatin	Echinacea does not appear to alter the AUC and clearance of oral midazolam, although the bioavailability may be increased. Clearance of intravenous midazolam may be modestly increased in patients taking <i>Echinacea</i> .
Cod liver oil	Ganfort	There are no interaction results for ganfort and cod liver oil
Hawthorne	Nitidipine	Limited evidence suggests that there may be additive blood pressure-lowering effects if hawthorn is taken with conventional antihypertensives, but the effects are small.

Visionace (lutein, carotenoids, myrtillus, flavonoid compounds)	Lansoprazole	The desired effect of betacarotene supplementation may be reduced in those taking proton pump inhibitors.
Evening primrose oil	Aspirin	Evening primrose oil can inhibit platelet aggregation and increase bleeding time. It has therefore been suggested that it may have additive effects with other antiplatelet drugs, but evidence of this is generally lacking.
Nytol (herbal)	Trimipramine	There are no interaction results for trimipramine and nytol herbal

* Potential interaction reports from Stockley's Herbal Medicine Interaction.

^[a] The number of patients exposed to the particular combination of HMPS /dietary supplement and prescription drug

^bGuidance about possible adverse effects, and/or some monitoring may be needed