

1 **Identifying patterns of general practitioner service utilisation and their relationship**
2 **with potentially preventable hospitalisations in people with diabetes: The utility of a**
3 **cluster analysis approach**

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27 **Abstract**

28 **Aims:** We aimed to characterise use of general practitioners (GP) simultaneously
29 across multiple attributes in people with diabetes and examine its impact on diabetes
30 related potentially preventable hospitalisations (PPHs).

31 **Methods:** Five-years of panel data from 40,625 adults with diabetes were sourced
32 from Western Australian administrative health records. Cluster analysis (CA) was
33 used to group individuals with similar patterns of GP utilisation characterised by
34 frequency and recency of services. The relationship between GP utilisation cluster
35 and the risk of PPHs was examined using multivariable random-effects negative
36 binomial regression.

37 **Results:** CA categorised GP utilisation into three clusters: moderate; high and very
38 high usage, having distinct patient characteristics. After adjusting for potential
39 confounders, the rate of PPHs was significantly lower across all GP usage clusters
40 compared with those with no GP usage; IRR=0.67 (95%CI: 0.62-0.71) among the
41 moderate, IRR=0.70 (95%CI 0.66-0.73) high and IRR=0.76 (95%CI 0.72-0.80) very
42 high GP usage clusters.

43 **Conclusions:** Combination of temporal factors with measures of frequency of use of
44 GP services revealed patterns of primary health care utilisation associated with
45 different underlying patient characteristics. Incorporation of multiple attributes, that
46 go beyond frequency-based approaches may better characterise the complex
47 relationship between use of GP services and diabetes-related hospitalisation.

48 **Keywords:** Cluster analysis; primary health care; potentially preventable
49 hospitalisation; diabetes; data linkage.

50

51

52 **1. Introduction**

53 Diabetes is an increasing public health issue causing a substantial burden on health
54 care systems around the world [1]. In Europe, the number of people with diabetes
55 was nearly 60 million in 2013, and is estimated to increase to 70 million by the early
56 2030s [2]. Similarly, in the United States the prevalence of diabetes was estimated at
57 29.1 million in the national report in 2014 [3]. In Australia, a country of approximately
58 24 million people, the prevalence of diabetes was about 1.2 million in 2014-15 [4]
59 and is estimated to increase to 3.4 million by early 2030s [5]. The condition costs the
60 Australian Health system more than \$AU6.5 billion each year [5]. Diabetes is
61 considered an ambulatory care sensitive condition [5], and consequently enhancing
62 primary health care to better manage diabetes has been a major approach in the
63 health care system of Australia [5, 6].

64 The literature suggests that better primary health care delivery reduces the risk of
65 hospitalisations for ambulatory care sensitive conditions in general [7-9]. With
66 respect to diabetes, a recent systematic review indicated that regular primary care
67 was associated with reduced risk of hospitalisation [10]. However, other aspects
68 such as frequency of visits or access to primary health care show inconsistent
69 results [10].

70 In Australia, primary care services, mainly provided by general practitioners (GP),
71 are subsidised through a universal health insurance scheme, Medicare, on a fee-for-
72 service basis [6]. Dedicated financial incentives have been provided under Medicare
73 for GPs to provide comprehensive care for diabetes [6]. However, to our knowledge,
74 limited research has evaluated patterns of utilisation of primary health care services
75 for people with diabetes and their impact on health outcomes. Current studies are
76 limited to examining the utilisation of primary health care based on single indicators
77 such as frequency [6] or regularity of services used [11].

78 Since patterns of primary health care utilisation are likely to be complex, more
79 advanced approaches that account for multiple factors are required to more
80 accurately classify and discover meaningful patterns of primary health care utilisation
81 by people with diabetes. K-mean cluster analysis, a data-driven approach, is capable
82 of taking into account multiple dimensions simultaneously and is suitable for use with
83 large datasets [12]. The technique can classify individuals with similar characteristics
84 into homogeneous groups which can also maximise heterogeneity between groups

85 [12]. The technique has been applied to a variety of settings, for example, health
86 behaviour [13]; health psychology [14]; health care cost analysis [12] and genetic
87 classification [15].

88 Thus, our study aims to apply K-mean cluster analysis to identify GP utilisation
89 patterns using multiple attributes of GP usage among people with diabetes. We will
90 also examine the impact of identified GP utilisation patterns on the risk of potentially
91 preventable hospitalisations (PPHs). Understanding patterns of GP utilisation and
92 how they impact on health outcomes is useful for planning health care provision
93 targeted to encouraging particular patterns in utilisation and enhancing the
94 relationship between patients and their primary health care provider.

95 **2. Material and methods**

96 **2.1 Data sources**

97 The Western Australian (WA) linked data used for this study comprised whole-of-
98 population administrative health data linked at the individual level, for residents of
99 WA aged 18 years or older who were registered at any time on the WA Electoral
100 Roll [16]. The data included a complete set of WA Hospital Morbidity Data
101 System (HMDS) records; Medicare Benefit Scheme (MBS) claim records; WA
102 Electoral Roll (ER) records; and WA mortality records for each individual
103 subsequent to their first ever WA Electoral Roll record. Details of each dataset
104 have been described previously [17]. In brief, the datasets provide statutory
105 information on all hospitalisations (HMDS), claims for medical services out-of-
106 hospital including GP visits (MBS), dates individuals migrated in and out of WA or
107 changed address while living in WA (Electoral Roll) and date/cause of death.

108 **2.2 Study population**

109 Annual panel data from 1998/1999 to 2003/2004 were constructed consisting of
110 individuals with diabetes identified via HMDS or MBS data prior to the start of or
111 in the baseline financial year (1998/99). Diabetes mellitus was determined using
112 the International Classification of Disease (ICD), 9th edition-clinical modification
113 (ICD-9-CM) codes in HMDS records and MBS claims indicative of the presence
114 of diabetes as described elsewhere [17]. All individuals were observed annually
115 from the baseline year to 30 June 2004, last year living in WA or death
116 (whichever occurred first) for any change in GP utilisation, hospitalisations and
117 clinical and demographic characteristics. GP utilisation and demographic and

118 clinical characteristics were measured in the exposure year, and PPH outcomes
119 measured in the following year. Only individuals who were alive and resident in
120 WA for at least two consecutive years were included in the study. The couplet
121 design (ie. comprising pairs of years, the exposure year followed by an outcome
122 year) has been applied in recent publications [6, 17].

123 Ethical approval was provided by The University of Western Australia and Curtin
124 University Human Research Ethics Committees who exempted the study from
125 obtaining individual patient consent.

126 **2.3 Study outcome and predictors**

127 **2.3.1. Diabetes related potentially preventable hospitalisations**

128 The primary outcome measure was diabetes related potentially preventable
129 hospitalisations (PPH) during the following-up year of each couplet.
130 Hospitalisations were deemed PPHs based on either their principal diagnosis
131 being identified by the National Health Performance Framework [18] as a
132 diabetes related PPH or identification by Davis et al [19] as associated with
133 increased risk for people with diabetes. Principal diagnoses were captured using
134 ICD-9-CM and Australian Modification ICD codes 10th revision (ICD-10-AM)
135 codes included in the HMDS records (Appendix 1).

136 **2.3.2. Variables for GP usage clustering**

137 The goal of these cluster analyses was to identify patterns of GP service
138 utilisation among people with diabetes. Candidate variables included in the
139 cluster analyses were adapted from the customer relationship management
140 framework proposed by Hughes (2005) [20] that capture both level of usage and
141 strength of the relationship between patients acting as customers and GPs acting
142 as primary care providers. Three main components suggested from the
143 framework were Recency, Frequency and Monetary [20] which have been
144 applied to healthcare data previously [21]. Since healthcare costs for Australia
145 are covered by Medicare, with limited out of pocket payment from patients, the
146 monetary component was not considered in our analyses. Greater recency and
147 frequency are indicators of how well the relationship between patients with
148 diabetes acting in the role of a customer and primary health care provider (GP)
149 acting in the role of the service provider has been maintained [21].

150 In our study recency of GP usage consisted of three factors including: (i) the
151 average time interval between access of health care service capturing the overall
152 interaction between patients and GPs, (ii) the standard deviation from the
153 average time interval capturing the extent of consistency in service utilisation,
154 and (iii) the longest time interval between services capturing the extent that
155 patients were out of coverage of primary care. Since the mean and standard
156 deviation values may be driven by extreme values, two alternatives to the
157 recency variable group were also considered in the cluster analyses including (A)
158 mean time interval, mean absolute deviation from the mean and the longest time
159 interval and (B) median time interval, median absolute deviation from median,
160 and the longest time interval. The results of cluster analysis of the three groups of
161 variables were compared in table 1. The time interval was determined between
162 the date of a GP visit and the date of the previous health care service provided
163 either from a GP or hospitalisation.

164 Frequency of GP usage was defined as the number of GP visits in a financial
165 year. Those GP visits occurring within 14 days of the previous GP visit were
166 counted as one GP usage to minimise over counting GP service utilisation, as
167 those within 14 days of each other are likely to be associated with a single
168 episode of care, for example where people may need to return to a GP to receive
169 laboratory test results, rather than a subsequent discrete GP service as
170 discussion with our GP experts.

171 All indicators were measured within financial years. However, a three-year look-
172 back period was used, where necessary, to calculate the time interval between
173 the first GP service in that year and the previous service. Three years was found
174 to be the tie period that maximised capturing recency of GP utilisation for the
175 cohort. Individuals having only one GP visit within a financial year were included
176 in the cluster analysis if they had a previous health care service within the look-
177 back period to enable the calculation of recency of GP usage.

178 **2.3.3. Covariates**

179 For this study, a number of individual characteristics were included to control for
180 potential confounders in the relationship between GP usage cluster and PPHs.
181 Demographic characteristics included were age group (18-44, 45-59, 60-74 and
182 ≥75 years), gender, Indigenous status, quintile of the Census specific Socio-
183 Economic Indexes for Areas (SEIFA) Index of Relative Socioeconomic

184 Disadvantage [22] and quintile of accessibility to services [23]. Diabetes
185 complications were identified using ICD codes suggested by Young, Lin [24] and
186 classified into four groups (0, 1, 2 and 3 or more complications) according to our
187 previously published methods [17]. The number of comorbidities was summed
188 from a list of comorbidities suggested by Holman et al. [25], excluding conditions
189 classified as complications of diabetes. Regularity of GP visits was calculated as
190 $[1/(1+\text{variance})]$ [9], where variance is a variance of the time interval between GP
191 visits occurring within the financial year and classified into four quantiles. Number
192 of specialist visits, and non-diabetes related hospitalisation were calculated within
193 a financial year. Duration of diabetes was calculated in years.

194 **2.4 Statistical analyses**

195 Cluster analyses were conducted using different alternative combinations of
196 recency and frequency of GP usage among those with at least one GP visit in a
197 financial year. First, the values of the mean/median time interval, the standard
198 deviation/absolute deviation of mean/median time intervals, longest time interval
199 and frequency of GP visits were normalised by subtracting the minimum of each
200 value and dividing that difference by the range of all values [12]. K-mean cluster
201 analyses were then conducted on normalised values of recency and frequency of
202 GP visits. The K-mean cluster approach was preferred as it is less susceptible to
203 outliers in the data and is appropriate for use with large datasets [12]. The
204 number of clusters was indicated using Calinski-Harabasz stopping rules for the
205 options of 2 to 6 clusters, the large values of the Calinski-Harabasz pseudo-F
206 index indicated distinct clustering [26]. Characteristics of final GP usage clusters
207 were described using a box plot.

208 Both descriptive bivariate and multivariate analyses were performed. Descriptive
209 analyses were used to summarise characteristics of participants among no GP
210 usage and each GP usage cluster in the baseline year. The results were
211 presented as the mean and standard deviation (SD) for continuous variables and
212 percentage for categorical variables. Multivariate analyses were conducted using
213 random-effects negative binomial regression model (NB) for panel data and zero-
214 inflated negative binomial regression model (ZINB) with the inflated component
215 contained in the intercept only. The Bayes Information Criterion (BIC) and Akaike
216 Information Criterion (AIC) statistics were used to assess the fit of the model
217 where NB with random effects was the preferred model compared to ZINB. We

218 included Mundlak variables, defined as group-means of time-varying variables, to
219 relax the assumption in the random-effects estimator that observed covariates
220 were uncorrelated with the unobserved covariates [27, 28]. The group mean
221 variables used were number of specialist visits and non-diabetes related
222 hospitalisation. All analyses were conducted using STATA for Windows version
223 14.1.

224 **3. Results**

225 *Clustering results*

226 Table 1 presents summary results of cluster analyses with different groups of
227 recency variables. The candidate group included mean time interval, mean
228 absolute deviation from the mean, longest time to GP visit and frequency of GP
229 visits; alternative A group included mean, standard deviation, the longest time
230 interval and frequency; alternative B group included median, median absolute
231 deviation from median, the longest time interval to GP visit and frequency of GP
232 visits. Using the Calinski cluster stopping rule, all three groups identified three
233 clusters. Compared with the candidate group, the other alternative groups had
234 very high percentage of agreement in term of grouping subjects into a cluster with
235 99.3% in the alternative A group and 95.5% in the alternative B group. The
236 candidate group also had highest Calinski F index value. Thus, the results of the
237 candidate group were kept to present in this paper (Table 1). Figure 1 and Table
238 2 summarise the GP usage clusters from K-mean analyses. Three clusters were
239 identified, including 1) moderate GP usage with mean time interval of
240 approximately 10 months (296 days), standard deviation of about 4 months (115
241 days), the longest time interval of 14 months (404 days) and frequency of about
242 2 times a year; 2) high GP usage with mean time interval to a GP visits of 3
243 months (88 days), standard deviation of 1.5 months (48 days), the longest time
244 interval of 5 months (147 days) and frequency of 3.7 times a year; and 3) very
245 high usage with mean time interval of 1.5 months (40 days), deviation of 0.5
246 months (20 days), the longest time interval of 2 months (76 days) and frequency
247 of visit approximately 7.8 times a year.

248 *Characteristics of study population by GP usage cluster at the baseline year*

249 Basic demographic and clinical characteristics of the study population are
250 described in Table 3 by no GP usage and each GP usage cluster. The majority of
251 the study population had high (n=17 077, 42.0%) and very high (n=15 858,

252 39.0%) GP usage, were aged 45 years or older (86.2%), and were more likely to
253 be male (51%), non-indigenous (92.7%), moderate to least disadvantaged
254 (51.6%), and living in areas with moderate to high accessibility to services
255 (93.4%). Those with complications accounted for 43.3%% in the study
256 population, higher in very high GP usage cluster (51.5%). The average number of
257 comorbidities was 4.5 (SD3.6), the highest in those with very high GP usage
258 cluster (mean 5.6; SD 3.5), followed by high GP usage cluster (mean 4.1, SD
259 3.5), no GP usage cluster (mean 3.5; SD 4.4) and moderate GP usage cluster
260 (mean 3.2; SD 2.9). The average duration of diabetes was 6.4 (SD=4.3) years,
261 similar duration across GP usage clusters and the no GP usage group. None and
262 low regularity of GP visits were observed across GP usage clusters, except the
263 very high GP usage cluster. High numbers of hospitalisations were observed
264 among those with no GP usage (average of 3.4 admissions), followed by the very
265 high GP usage cluster (0.8 admissions), high GP usage cluster (0.7 admissions)
266 and moderate GP usage cluster (0.2 admissions).

267 Overall, the moderate GP usage cluster tended to be younger (25.1% aged 18-44
268 years, and 37.7% aged 45-60 years), male (62.6%), Indigenous (10.1%), live in
269 less accessible areas (25.7%), compared with both the high and very GP usage
270 cluster (Table 3). The moderate GP usage cluster was less likely to have
271 complications (27.2%); had a lower number of comorbidities (3.2 (SD 2.9)); was
272 less likely to have regular GP visits (20.5%) and had a lower number of
273 hospitalisation (0.2; SD 0.8) compared with both high and very GP usage clusters
274 The no GP usage group was quite comparable to other GP usage clusters in
275 term of age, gender, complications and comorbidity distribution. However, the no
276 GP usage group had a higher proportion of individuals who were indigenous
277 (23.7%), in the highest disadvantage SEIFA quintiles (31.1%) and resided in very
278 remote areas (20.1%).

279

280 *Association between GP usage and the risk of hospitalisations*

281 The preferred model was the panel negative binomial regression model based on
282 information criterion (AIC and BIC). The results show that GP usage across all
283 clusters had a protective effect against the risk of PPH in the following year after
284 adjusting for all covariates. However, the greatest protective effect was observed
285 for individuals in the moderate GP usage cluster (IRR=0.67 (95%CI: 0.62-0.71).

286 The average adjusted predictions indicate that on average 0.25 PPHs per year
287 (95%CI: 0.24-0.27) can be expected for those in the moderate GP cluster; 0.26
288 per year (95%CI 0.259-0.27) for those in the high GP usage cluster and 0.29 per
289 year (95%CI: 0.28-0.30) for those in the very high GP usage cluster, while those
290 with no GP usage are estimated to have on average 0.38 hospitalisations per
291 year (95%CI: 0.36-0.40) (Figure 2).

292

293 **4. Discussion**

294 This study aimed to reveal the latent pattern of GP contact using K-mean cluster
295 analysis, a novel statistical technique, which overcomes many of the limitations
296 associated with current studies by examining GP service use simultaneously
297 across multiple attributes. Importantly we were able to include time intervals
298 between service utilisations including average time interval, deviation of the time
299 intervals and the longest time interval in assessing the patterns of GP service
300 use which enhance the classification accuracy.

301 The rationale behind our exploration of incorporating multiple attributes to
302 categorise GP use is our hypothesis that using frequency or regularity of GP
303 contact alone may be too simplistic, since individuals that have the same number
304 of visits or the same regularity in a year may have differences in the temporal
305 distribution of visits. Shorter time intervals between services in combination with
306 more regular provision may reflect “proactive care” and the strengthening of the
307 relationship between patients and their GP. In turn, proactive care may allow the
308 opportunity for continuous improvement in self-management skills and health
309 literacy which may assist in the prevention and early treatment strategies in the
310 primary care setting [6, 29]. The characterisation of GP utilisation based on
311 multiple domains of GP use has not to our knowledge been previously reported
312 and, we argue represents an advance on current single domain methods.

313 In our study, although the no GP usage group was comparable to other GP
314 usage clusters in term of age and gender and disease severity, the group
315 comprised higher proportion of disadvantage population (Indigenous status,
316 highest disadvantage SEIFA and very remote). These findings highlight the
317 existence of inequity in access of primary care for people with diabetes in
318 particular sub-populations which have been previously reported in the literature

319 [30, 31] .The majority of individuals with diabetes were categorised in high or
320 very high GP usage clusters. Those in high and very high GP usage clusters had
321 high and very high recency and frequency of GP usage, respectively while those
322 in the moderate GP usage cluster had both lower recency and frequency of
323 contact. The clinical characteristics of each cluster differed significantly with
324 those in the high or very high GP usage clusters more likely to have a higher
325 number of complications and comorbidities compared with the moderate GP
326 usage cluster. These results were in line with literature that showed higher health
327 care service utilisation was observed among diabetes with multiple comorbidities
328 and complications [32-34]. Thus, the multidimensional GP usage clusters
329 identified in our study may be an indicator of patients' clinical characteristics
330 which is driving their health care needs. This represents an improvement on
331 other more simplistic measures such as frequency that do not correlate well with
332 health outcomes [6, 10].

333 The literature does not show a consistent relationship between the level of
334 primary health care and the risk of hospitalisation [7, 10]. While Comino et al.
335 found that higher number of GP visits increased the risk of hospitalisation [6],
336 other authors found an inverse relationship between the frequency of GP visits
337 and hospitalisation [35]. Discordant results in the literature may be due to the
338 complexity of the mechanism in the relationship between primary health care
339 and hospitalisation, which may not be adequately captured by the number of GP
340 visits [6]. Thus, use of a more complex measure of GP use, such as that
341 developed in our study which incorporates several dimensions may be better
342 suited to understand the risk of hospitalisation and help predict and contain the
343 costs of healthcare for diabetes.

344 Our findings support the hypotheses that GP contact reduces the risk of
345 hospitalisation. However, the effect was not linear for each additional level of GP
346 usage, with the highest effect observed among those with moderate GP usage
347 cluster. This may be explained by characteristics of GP usage cluster, those with
348 moderate usage were likely to be younger, have fewer complications and
349 comorbidities than those with high and very high GP usage. The results were
350 also supported by the health demand model of Grossman where health is
351 considered as a durable capital stock that depreciates with age and can be

352 increased through investment in healthcare [36]. Thus, a finite lifetime increase
353 in the depreciation rate of health may lead to an increase in demand for both
354 preventive care and curative care [36, 37]. However, if primary health care can
355 provide early treatment and prevention of illness, it would still be a substitute for
356 hospital care in some instances [37].

357 *Strengths and limitations of the study*

358 The major strength of our study is that it was based on a large set of linked
359 administrative data at the individual level that encompassed the whole-
360 population and a comprehensive range of health care services. The linked
361 whole-of-population data allowed us to assess changes in both exposure and
362 outcomes at the individual level over the follow-up period. The panel data
363 structure contained information on both within and between individual variations
364 enabling us to control for the effect of unobserved covariates [38]. Our study also
365 applied a novel advanced analytic approach, cluster analysis, and customer
366 relationship management framework to reveal previously hidden patterns of
367 primary health care utilisation. These approaches allowed us to examine primary
368 health care utilisation across multiple attributes simultaneously, and thus
369 characterise a measure of GP utilisation that may facilitate a better
370 understanding of the influence of primary health care in reducing the risk of
371 hospitalisations among people with diabetes.

372 Our study has some limitations. Comorbidity was accessed by a simple count of
373 conditions which may not well capture actual health care needs although the
374 measure is frequently used in the literature [6, 34, 39]. The analyses were limited
375 to Australian citizens in one Australian State, due to the reliance on the WA
376 Electoral Roll, and those with a previous diagnosis of diabetes captured by our
377 data. Thus, the result may not be fully generalizable to all individuals living with
378 diabetes, since the Electoral Roll is known to under-represent some groups such
379 as Indigenous Australians and those aged under 21 years of age [40]. However,
380 the use of longitudinal Electoral Roll data provided the ability to accurately
381 capture person-time at risk, due to capturing movement in and out of the state
382 [40]. Limiting the study to a single Australian State is unlikely to have significantly
383 influenced the findings, since Australia has a single public health system,
384 Medicare. Similarly, our reliance on linked administrative health data to identify

385 those diagnosed with diabetes limited the study to those who have previously
386 accessed health services pathognomonic of diabetes and thus people living with
387 diabetes who have never accessed diabetes-related health services are not
388 represented. Individuals not included in our data are likely to be the lower
389 severity patients who are less likely to need hospital care. These limitations are
390 common and well-known in administrative datasets and, because of the features
391 of the excluded patients, are likely to have limited effect on our examination of
392 the pattern of primary care utilisation and the relationship between the patterns
393 of utilisation on the risk of hospitalisation in previously diagnosed diabetes.

394 Through combining both temporal factors with measures of frequency of use of
395 GP services our study revealed a latent pattern of primary health care utilisation.
396 Incorporation of multiple attributes that go beyond a simplistic frequency-based
397 approach may better characterise the complex relationship between use of GP
398 services and diabetes-related hospitalisation. The study has demonstrated the
399 ability of cluster analyses to provide a systematic formalised approach for
400 exploring complex patterns of health service utilisation in large administrative
401 datasets. Application the cluster analysis approach to other chronic conditions
402 would be useful for accurate understanding patterns of service utilisation. Future
403 studies should further examine temporal factors in the provision of primary health
404 care and evaluate what combination of time between visits, regularity and
405 frequency of access to primary care would best improve health outcome and
406 contain costs.

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413

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515 **Results: Figures**

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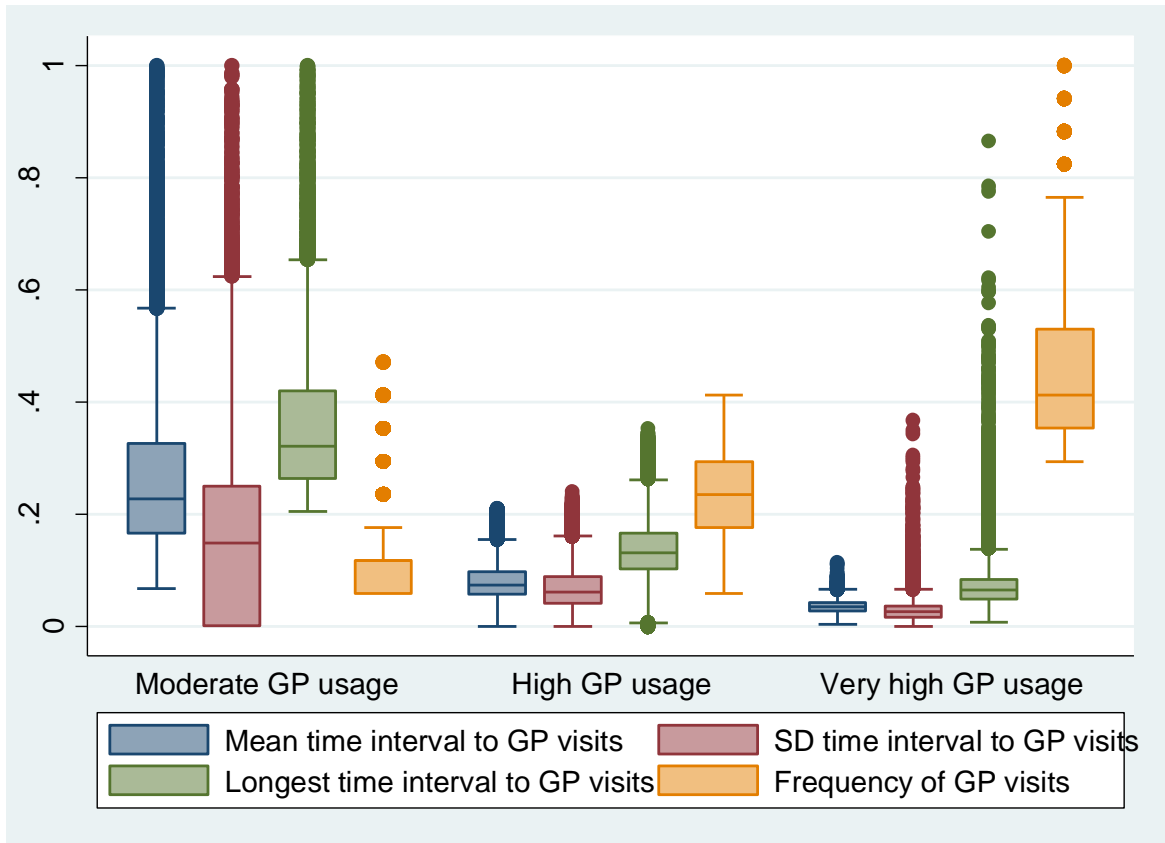


Figure 1. GP usage by clusters

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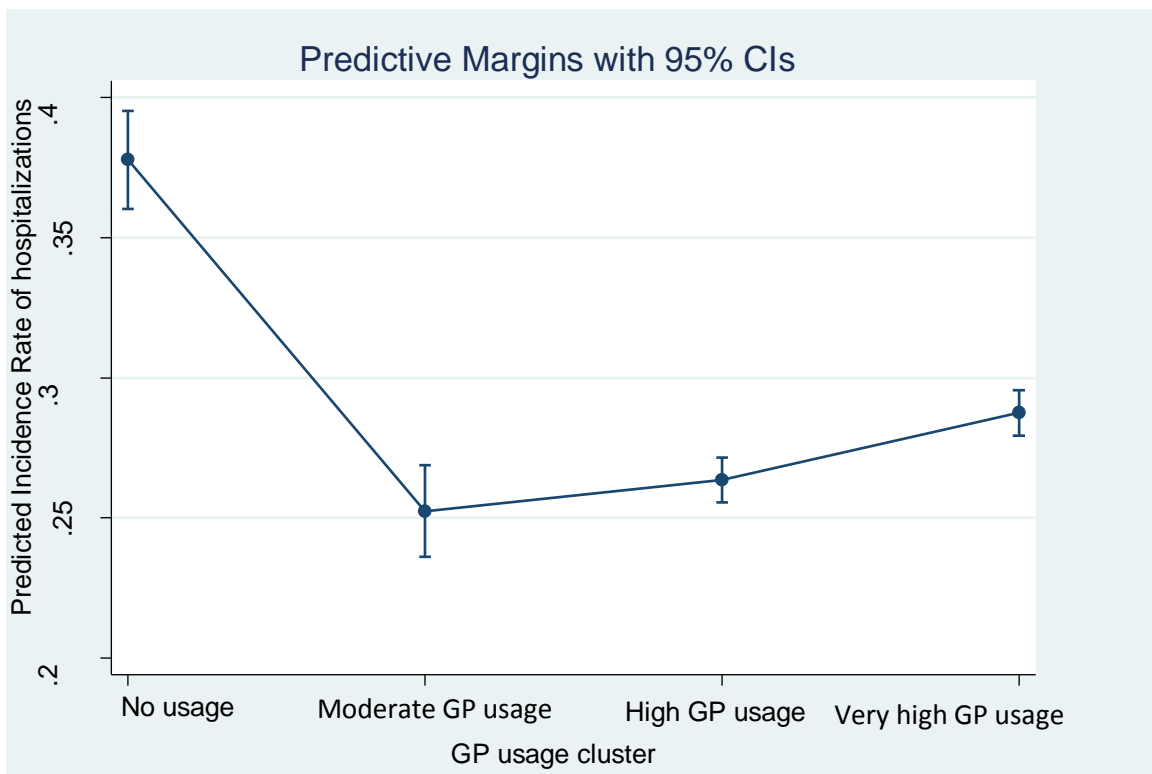


Figure 2. Predictive margins the incident rate of diabetes related PPH

525 **Results: tables**

526 *Table 1 Cluster analysis outputs with different groups of recency variables*

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	Group of Indicators used in K-mean cluster		
	Candidate group	Alternative A group	Alternative B group
Mean	✓	✓	
Median			✓
Mean absolute deviation from the mean		✓	
Median absolute deviation from median			✓
standard deviation	✓		
The longest time to GP visit	✓	✓	✓
Frequency of GP visits	✓	✓	✓
Cluster stopping (Caliński rule)	133805	132616	129095
Number of clusters	3	3	3
% of agreement vs. group 1 (Kappa values)	-	99.3%	95.5%

528

529 *Table 2 GP usage clusters summary*

Clusters	Mean (days)	SD (days)	The longest (days)	Frequency of GP visits
Moderate usage				
Min	75	0	225	1
Mean	296.8412	115.0688	404.0527	1.919529
Max	1093	744.5834	1095	8
High usage				
Min	1	0	1	1
Mean	88.19658	48.81665	147.0608	3.716618
Max	230	178.1975	387	7
Very high usage				
Min	5.2	0	9	5
Mean	39.71341	20.81995	76.12468	7.819856
Max	124.75	273.0432	947	17

530

531

Characteristics	No GP usage	Moderate GP usage	High GP usage	Very high GP usage
	(N, (%))	(N, (%))	(N, (%))	(N, (%))
N (%)	4 198 (10.3)	3 492 (8.6)	17 077 (42.0)	15 858 (39.0)
Age group (years)				
18-44	781 (18.6)	877 (25.1)	2,668 (15.6)	1178 (7.4)
45-59	1059 (25.2)	1316 (37.7)	5,649 (33.1)	3543 (22.3)
60-74	1183 (28.2)	1,016 (29.1)	6,655 (38.9)	7465 (47.1)
≥75	1175 (28.0)	283 (8.1)	2,105 (12.3)	3672 (23.2)
Gender				
Female	1679 (40.0)	1307 (37.4)	7,912(46.3)	9002 (56.8)
Male	2519 (60.0)	2185 (62.6)	9,165 (53.7)	6856 (43.2)
Indigenous status				
No	3084 (76.3)	2911 (89.8)	15,197 (93.8)	14978 (96.5)
Yes	961 (23.7)	329 (10.1)	1,003 (6.2)	549 (3.5)
SEIFA				
Highest Disadvantage	1285 (31.4)	631 (18.4)	23,240 (19.2)	3435 (21.8)
High disadvantaged	1037 (25.3)	918 (26.7)	4,797 (28.4)	4558 (28.9)
Moderate disadvantage	573 (14.0)	593 (17.3)	2,381 (14.1)	2185(13.8)
Less disadvantage	544 (13.5)	561 (16.3)	2,754 (16.3)	2416(15.3)
Least disadvantage	645 (15.7)	728 (21.2)	3,691 (21.8)	3158 (20.0)
Accessibility				
Very remote	825 (20.1)	251 (7.3)	611 (3.6)	79 (1.2)
Remote	172 (4.0)	90 (2.6)	355 (2.1)	184 (1.1)
Moderate	268 (6.5)	265 (7.7)	946 (5.6)	659 (4.2)
Accessible	210 (5.1)	273 (7.9)	1,027 (6.1)	695 (4.4)
Highly accessible	2619 (63.9)	2,552 (74.3)	13,926 (82.6)	14036 (89.1)
Complication severity level				
No complication	1957 (46.6)	2,543 (72.8)	10,845 (63.5)	7694 (48.5)
1 complication	746 (17.8)	385 (11.0)	2,372 (13.9)	2638 (16.6)
2 complications	577 (13.7)	322 (9.2)	1,804 (10.5)	2266 (14.3)
3+ complications	918 (21.9)	242 (6.9)	2,056 (12.0)	3260 (20.6)
Number of comorbidity				
Mean (SD)	3.5 (4.4)	3.2 (2.9)	4.1 (3.4)	5.6 (3.5)
Duration of diabetes (years)				
Mean (SD);	6.7 (4.4)	6.3 (4.2)	6.1 (4.2)	6.5 (4.4)
Regularity quantiles				
No regularity	4,198 (100.0)	2,776(79.5)	3,315 (19.4)	0
Quantile 1		716 (20.5)	6,684 (39.1)	287 (1.8)
Quantile 2			4,719 (27.6)	2,972 (18.7)
Quantile 3			1,497 (8.8)	5,917 (37.3)
Quantile 4			862 (5.0)	6,682 (42.1)
Diabetes related PPH				
Mean (SD)	2.5 (17.5)	0.07 (0.38)	0.25 (2.6)	0.25 (1.02)

Table 4 Association of GP usage pattern and potentially preventable hospitalisation with and without adjustment for other covariates (results from random effects negative binomial regression)

	Multivariate NB		Adjusted multivariate NB		ZINB	
	IRR	(95%CI)	IRR	(95%CI)	IRR	(95%CI)
GP cluster usage						
No usage	1	(1; 1)	1	(1; 1)	1	(1; 1)
Moderate usage	0.62***	(0.57; 0.66)	0.67***	(0.62; 0.72)	0.41***	(0.33; 0.50)
High usage	0.67***	(0.64; 0.71)	0.70***	(0.66; 0.73)	0.40***	(0.35; 0.46)
Very high usage	0.76***	(0.72; 0.79)	0.76***	(0.72; 0.80)	0.39***	(0.34; 0.45)
Gender						
Males vs. females	1.06***	(1.03; 1.10)	1.07***	(1.04; 1.11)	1.24***	(1.13; 1.36)
Age (years)						
18/44	1	(1; 1)	1	(1; 1)	1	(1; 1)
45/59	1.20***	(1.12; 1.28)	1.21***	(1.14; 1.29)	1.10	(0.91; 1.32)
60/74	1.74***	(1.64; 1.86)	1.73***	(1.62; 1.84)	1.44***	(1.20; 1.73)
75+	2.30***	(2.15; 2.46)	2.31***	(2.16; 2.47)	1.42***	(1.18; 1.71)
Indigenous status						
Yes vs. No	1.47***	(1.37; 1.59)	1.50***	(1.39; 1.61)	2.18***	(1.79; 2.67)
SEIFA						
Highest Disadvantage	1	(1; 1)	1	(1; 1)	1	(1; 1)
High disadvantaged	0.95*	(0.91; 1.00)	0.95*	(0.91; 0.99)	0.96	(0.84; 1.09)
Moderate disadvantage	0.95	(0.90; 1.00)	0.94*	(0.89; 0.99)	0.86*	(0.76; 0.97)
Less disadvantage	0.98	(0.93; 1.03)	0.97	(0.92; 1.02)	0.95	(0.82; 1.10)
Least disadvantage	0.93**	(0.88; 0.98)	0.90***	(0.86; 0.95)	0.94	(0.81; 1.09)
Accessibility						
Very remote	1	(1; 1)	1	(1; 1)	1	(1; 1)
Remote	1.00	(0.87; 1.13)	1.00	(0.88; 1.13)	0.76*	(0.59; 0.96)
Moderate	0.97	(0.88; 1.08)	0.98	(0.88; 1.08)	0.84	(0.64; 1.09)
Accessible	0.92	(0.83; 1.03)	0.92	(0.82; 1.02)	0.73*	(0.57; 0.95)
Highly accessible	0.89*	(0.82; 0.98)	0.90*	(0.83; 0.99)	0.97	(0.78; 1.21)
Duration of diabetes (years)	1.03***	(1.03; 1.04)	1.04***	(1.03; 1.04)	1.05***	(1.04; 1.06)
Complication severity level						
No complication	1	(1; 1)	1	(1; 1)	1	(1; 1)
1 complication	1.33***	(1.27; 1.40)	1.27***	(1.21; 1.33)	1.05	(0.94; 1.18)
2 complications	1.68***	(1.60; 1.77)	1.58***	(1.51; 1.66)	1.57***	(1.37; 1.80)
3+ complications	2.12***	(2.02; 2.22)	1.90***	(1.81; 2.00)	2.72***	(2.34; 3.15)
Number of comorbidities						
Number of specialist services	1.07***	(1.06; 1.07)	1.04***	(1.03; 1.04)	1.07***	(1.05; 1.09)
Non-diabetes related hospitalisation	1.01***	(1.01; 1.01)	0.99***	(0.98; 0.99)	0.97***	(0.96; 0.98)
Diabetes related hospitalisation lag1	1.05***	(1.02; 1.09)	0.99	(0.96; 1.02)	0.99	(0.90; 1.10)
Diabetes related hospitalisation baseline			1.36***	(1.31; 1.40)	4.65***	(3.94; 5.49)
Group mean number of specialist visits			1.11***	(1.07; 1.14)	1.14*	(1.02; 1.27)
Group mean non-diabetes related hospitalisations			1.04***	(1.04; 1.05)	1.06***	(1.05; 1.08)
			1.60***	(1.50; 1.72)	1.89***	(1.52; 2.36)
AIC	191782.6		190686.5		202182.5	
BIC	192075.6		191019.9		202515.9	

Exponentiated coefficients

="* p<0.05

** p<0.01

*** p<0.001"

Table 5 Margin incident rate of diabetes related PPH

GP usage	Incidence rate	95% CI	
No GP usage	0.38	0.36	0.40
Moderate GP usage	0.25	0.24	0.27
High GP usage	0.26	0.26	0.27
Very high GP usage	0.29	0.28	0.30