

**Neuropsychiatric Symptoms as Predictors of Falls in Long-Term Care Residents with
Cognitive Impairment**

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26 **Abstract**

27 *Objectives:* Falls and neuropsychiatric symptoms (NPS) are common among long-term care
28 residents with cognitive impairment. Despite the high prevalence of both falls and NPS, little is
29 known about their association. The aim of our study was to explore how NPS, particularly the
30 severity of NPS and specific NPS subgroups, are associated with falls and how psychotropics
31 modify this association.

32 *Design:* Longitudinal cohort study.

33 *Setting and Participants:* 532 long-term care residents aged 65 years or over in Helsinki, Finland.

34 *Methods:* NPS were measured with Neuropsychiatric Inventory (NPI) at baseline. Participants were
35 grouped into three groups: no significant NPS (NPI points 0-3), low NPS burden (NPI 4-12) and
36 high NPS burden (NPI>12). The number of falls, injuries, fractures, and hospitalizations were
37 collected from medical records over 12 months following baseline assessment.

38 *Results:* Altogether 606 falls occurred during the follow-up year. The falls led to 121 injuries, 42
39 hospitalizations, and 20 fractures. Falls and injuries increased significantly with NPS burden
40 ($p<0.001$): 330 falls in the high NPS group ($n=184$), 188 falls in the low NPS group ($n=181$) and 88
41 falls in the no significant NPS group ($n=167$). The risk of falling showed a curvilinear association
42 with NPI total score. Of NPS subgroups, psychosis and hyperactivity were associated with a higher
43 incidence rate ratio of falls, whereas apathy had a protective association even after adjustment for
44 age, sex and mobility. Affective symptoms were not associated with falls. Psychotropics did not
45 modify the association between NPS burden and falls.

46 *Conclusions and Implications:* The results of this study show that NPS, especially NPS severity,
47 may predict falls and fall-related negative consequences. Severity of NPS should be taken into
48 account when assessing fall risk in long-term care residents with cognitive impairment.

49 **Keywords:** falls– neuropsychiatric symptoms – cognitive impairment – Neuropsychiatric Inventory
50 – long-term care

51 **Introduction**

52 Falls and fall-related negative consequences among long-term care residents with cognitive
53 impairment are common.¹⁻⁴ An estimated 37-65% of older people with cognitive impairment or
54 dementia fall annually.⁵⁻⁶ The risk factors for falls are multiple and seem to vary between
55 community- and institution-dwelling older adults with cognitive impairment.^{3, 6-7} Impaired mobility,
56 use of psychotropic drugs, anxiety, depression and orthostatic hypotension have been shown to
57 increase fall risk.^{4, 7-12}

58 Neuropsychiatric symptoms (NPS), also called behavioral and psychological symptoms of dementia
59 (BPSD), are known to be highly common in cognitive impairment, especially in long-term care
60 settings. The prevalence of NPS in long-term care has been estimated to be as high as 82-92%.¹³⁻¹⁴
61 NPS include such symptoms as agitation, apathy, anxiety, aberrant motor behavior, delusions,
62 dysphoria, disinhibition, euphoria, hallucinations, and irritability.¹⁵ Cluster analyses have identified
63 four NPS subgroups: hyperactivity, psychosis, affective symptoms, and apathy.¹⁶

64 Despite the high prevalence of both NPS and falls among older adults with cognitive impairment in
65 long-term care, little is known about the association between NPS and falls. A few studies have
66 suggested NPS to be an independent risk factor for falls.¹⁷⁻²² However, it is not known whether
67 severity of NPS has an impact on fall rate. Thus, the aim of our study was to explore how NPS, and
68 more specifically the severity of NPS, are associated with falls and their consequences. Another aim
69 was to determine how specific NPS subgroups impact the incidence of falls.

70 **Methods**

71 *Study participants*

72 Participants were recruited to this longitudinal cohort study from institutional settings in Helsinki in
73 2017. From a random sample from 18 nursing homes, 544 volunteer residents were recruited to this
74 study. Participants' baseline assessment occurred between February 2018 and August 2018. All

75 participants who completed the Neuropsychiatric Inventory (NPI) at baseline (n=532) were
76 included in the study. The participants were followed for 12 months or until death.

77

78 *Measures*

79 Study nurses were trained to perform the assessments. One of the researchers (HMR) participated in
80 and supported the nurses in the baseline assessments and data collection. Data on demographic
81 factors such as age, sex, and diagnoses, were collected from medical records. The study nurses
82 calculated the Charlson Comorbidity Index²³ to assess each resident's burden of comorbidity and
83 the Barthel Index²⁴ for functional evaluation. Mobility was assessed by the 15D questionnaire²⁵
84 item on mobility and categorized into one of the following: 1) "able to walk without help outdoors
85 or indoors", 2) "able to walk indoors only with help from others", or 3) "completely bed-ridden and
86 unable to move about." Mini Mental State Examination (MMSE)²⁶ and Clinical Dementia Rating
87 (CDR)²⁷ were performed to assess the severity of cognitive impairment. Presence of vision and
88 hearing deficits were assessed by 15D questionnaire²⁵ items on vision and hearing. Vision was
89 categorized into either 1) "able to read papers and/or TV text with or without glasses" or 2) "not
90 able to read papers or TV text either with glasses or without. Hearing was categorized into either 1)
91 "able to hear speech with or without a hearing aid with normal or louder than normal voice or 2)
92 "able to hear even loud voices poorly or deaf".

93 Data on medication use were retrieved from medical records on the assessment day. All
94 medications were classified using the Anatomical Therapeutic Chemical (ATC) classification
95 system.²⁸ Psychotropic medications included antipsychotics (N05A), antidepressants (N06A),
96 anxiolytics (N05B), and hypnotics and sedatives (N05C). The use of Alzheimer medication (N06D)
97 included cholinesterase inhibitors (N06DA) and/or memantine (N06DX01). Only regularly used
98 medications were considered. Medication use was considered regular if there was a documented
99 regular sequence of administration.

100 To evaluate NPS, study nurses interviewed care staff from the long-term care units using the NPI.²⁹
101 The NPI was chosen instead of NPI-NH, because the nursing home version has not been translated
102 to Finnish language. The content of the questions of the NPI and NPI-NH are identical. NPI
103 includes 10 different NPS (agitation, apathy, anxiety, aberrant motor behavior, delusions,
104 dysphoria, disinhibition, euphoria, hallucinations, irritability). For each symptom, the severity is
105 multiplied by the frequency, and the sum score provides the total NPI score (range 0 to 120).
106 Subgroups of “Psychosis” (delusion, hallucinations), “Hyperactivity” (agitation, euphoria,
107 disinhibition, irritability, aberrant motor behavior), “Affective symptoms” (depression, anxiety),
108 and “Apathy” (apathy) were calculated separately, as earlier described¹⁶. We grouped the residents
109 according to the total score on NPI into three groups: no significant NPS (NPI 0-3), low NPS
110 burden (NPI 4-12), and high NPS burden (NPI >12). According to previous studies, a score >3 is
111 taken to indicate the presence of clinically relevant symptoms.²⁹ The cut-off point of 12 was chosen
112 as it was the median. In previous studies a total NPI score greater than 11 points arising from at
113 least three domains has been considered to indicate marked neuropsychiatric symptoms.³⁰ After
114 baseline assessment, data regarding falls (number of falls, injuries, fractures, and hospitalizations)
115 were collected from medical records during the 12-month-follow-up. Mortality was retrieved from
116 central records.

117

118 *Statistics*

119

120 Data are presented as means with standard deviation (SD) or as counts with percentages. The
121 statistical significance for the unadjusted hypothesis of linearity across categories (tertiles) of NPI
122 total score and characteristics of study participants was evaluated using the Cochran-Armitage test
123 for trend, analysis of variance (ANOVA), and logistic (ordinal) models with an appropriate contrast.
124 A bootstrap method was used when the theoretical distribution of the test statistics was unknown or
125 in case of violation of assumptions (e.g. non-normality). We used the Kaplan-Meier method to

126 construct estimated mortality. Cox proportional Hazard Model was used to estimate age, sex and
127 mobility adjusted risk (HR) for mortality between the groups. The number and incidence rate of
128 falls were calculated assuming a Poisson distribution. Adjusted incidence rate and incidence rate
129 ratio (IRR) were calculated using a Poisson regression model that included sex, age and mobility as
130 covariates. Multivariate Poisson regression models with forward stepwise was used to investigate
131 factors related to incidence of falls. Variables significant at the $P < 0.10$ level in unadjusted analyses
132 were included into the model. Multivariate Poisson regression was tested using goodness of fit of
133 the model, and the assumption of overdispersion in the Poisson model was tested using the
134 Lagrange multiplier test. A possible non-linear relationship between all falls and the NPI total score
135 was assessed by using a 3-knot-restricted cubic (placed according to Harrell's recommended
136 percentiles) spline Poisson regression model. The normality of variables was evaluated graphically
137 and using the Shapiro–Wilk W test. Stata 16.0 (StataCorp LP, College Station, TX, USA) was used
138 for the analysis.

139

140 *Statement of ethics*

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142 The study protocol was approved by the Ethics Committee [REDACTED] Written
143 informed consent was obtained from each participant and in case of significant cognitive decline
144 (CDR 2 or 3) from their closest proxy.

145

146 **Results**

147 The three NPI groups were similar in baseline demographic characteristics such as age, sex and
148 Charlson Comorbidity Index (Table 1). Residents' mean age was 85 years, 80% were women and
149 the mean number of comorbidities according to the Charlson Comorbidity Index was 2.1.

150 Significant differences between the groups were detected in functional capacity according to the

151 Barthel Index and in mobility. The residents with the highest NPS burden were the most mobile and
152 had better functioning than residents with no significant NPS ($p < 0.001$). Two of three residents
153 suffered from severe cognitive impairment (CDR 3) and the mean MMSE was low, 6.8. No
154 significant differences existed between groups in severity of cognitive impairment according to
155 MMSE ($p = 0.89$) or CDR rating ($p = 0.35$).

156 The NPI groups differed significantly in psychotropic medication use ($p < 0.001$). Mean number of
157 psychotropic medications in the high NPS burden group was 2.3, compared with 1.8 in the group
158 with no significant NPS. The proportion of residents taking any psychotropic medication was very
159 high, 87%. Residents with high NPS burden were also administered more often Alzheimer's
160 medication ($p = 0.041$) and had a higher number of total medications 8.8, compared with 7.9 in the
161 group with no significant NPS ($p = 0.031$). The most common NPS subgroup was hyperactivity in all
162 NPI groups.

163

164 **Mortality and incidence of falls according to NPI total score during follow-up**

165 Total follow-up time was 446.8 person-years, with the mean time being 0.84 (range 0.01 – 1.00)
166 years per person. During the one-year follow-up the mortality was 28.7% in the group with no
167 significant NPS, 33.2% in the low NPS burden group, and 33.7% in the high NPS burden group
168 ($p = 0.56$). When the NPI 0-3 group was used as reference age, sex and mobility adjusted HR for
169 mortality in NPI 4-12 group was 1.08 (95% CI 0.73 – 1.60) and in NPI >12 group 1.19 (95% CI
170 0.80 – 1.79).

171 Altogether 606 falls occurred during the follow-up: 330 in the high NPS burden group, 188 in the
172 low NPS burden group and 88 in the no NPS group (Table 2). Of 532 residents, one-third fell at
173 least once (94 residents). Severity of NPS measured by NPI total score had a curvilinear association
174 with the incidence rate of falls per person years (Figure 1). Using the no significant NPS group as a
175 reference, the low NPS burden group had an IRR per SD for falls of 1.64 (95% CI 1.27 – 2.12,

176 adjusted for age, sex and mobility), whereas in the high NPS burden group the IRR per SD was 2.43
177 (95% CI 1.91 – 3.08, adjusted for age, sex and mobility) (p for linearity < 0.001).

178 The associations between NPS subgroups and the IRRs of falls and fall-related negative
179 consequences are presented in Figure 2. Psychosis and Hyperactivity subgroups were associated
180 with a higher IRR of falls and injuries, whereas Apathy showed a protective association against falls
181 but not injuries. Affective symptoms did not predict falls nor injuries. Psychosis, Hyperactivity and
182 Affective symptom subgroups were associated with a higher IRR of hospitalizations, whereas
183 Apathy was not. None of the subgroups predicted fractures (Figure 2).

184 Falls and fall-related negative consequences during the 12-month follow-up

185 Of 606 falls, 121 led to injuries, 42 to injuries needing hospitalization, and 20 to fractures. Falls and
186 injuries increased significantly with NPS burden ($p < 0.001$). Residents with a higher NPI total score
187 were also more often hospitalized for their falls than residents with no significant NPS or with a low
188 NPS burden ($p = 0.002$). The number of fractures increased in the higher NPI groups but it did not
189 reach statistical significance ($p = 0.16$) (Table 2).

190 In a multivariate poisson regression analysis a higher NPI level (“NPI 4-12” IRR 1.72, 95% CI 1.33
191 to 2.23; “NPI >12” IRR 2.58, 95% CI 2.03 to 3.29) and male gender (IRR 1.80, 95% CI 1.51 to
192 2.16) were associated with a higher incidence of falls. Worse mobility (“able to walk only with
193 help” IRR 0.57, 95% CI 0.47 to 0.69; “bed-ridden” IRR 0.10, 95% CI 0.07 to 0.15), age (IRR 0.98,
194 95% CI 0.97 to 0.99) and the use of psychotropic medication (IRR 0.88, 95% CI 0.83 to 0.94) were
195 associated with a lower incidence of falls. Alzheimer’s medication (IRR 1.12, 95% CI 0.94 to 1.33)
196 and hearing deficits (IRR 0.76, 95% CI 0.45 to 1.26) were not associated with incidence of falls
197 (Table 3).

198 Finally, we stratified the residents according to their psychotropic use to see how psychotropics
199 modified the association between NPS severity and incidence rate of falls. NPI level was associated
200 with incidence rate of falls per person-years ($p < 0.001$ for NPI level), whereas psychotropic drug use
201 did not have a significant association ($p = 0.94$ for psychotropic), and no interaction existed ($p = 0.57$
202 for interaction) (Figure 3).

203 **Discussion**

204 NPI total score of long-term care residents with cognitive impairment showed a curvilinear
205 association with the incidence rate of falls, indicating that severity of NPS is associated with risk of
206 falls. Even after adjustments, IRR per SD in the high NPS burden group was nearly 2.5-fold that in
207 the no significant NPS group. Psychotropic drug use did not modify this association. Another
208 important finding was that, of all the NPS subgroups, specifically Psychosis and Hyperactivity had
209 the highest association with fall rate, while Apathy seemed to have a protective association.

210 The results of this study are in line with the few previous reports examining the association between
211 NPS and falls.¹⁷⁻²² According to our study, the fall risk related to NPS seems to particularly arise
212 from hyperactivity and psychotic symptoms. A population-based study from Sweden in 2005 also
213 found that having hyperactive symptoms was one of the factors most strongly associated with falls.¹
214 The results from earlier studies regarding wandering are contradictory. A systematic review from
215 2013 found wandering to be protective against falls³, whereas more recent studies have suggested
216 that wandering increases the risk for falls.^{22, 31} In our study, affective symptoms and apathy were
217 not associated with an increased fall risk. This could be due to less day-time activity offering less
218 opportunities for falling.

219 The findings from previous studies indicate that both NPS and falls increase with the severity of
220 cognitive impairment.³²⁻³⁵ In our study, there was no difference in the severity of cognitive
221 impairment between the three NPI groups measured by MMSE or CDR. This could be due to the

222 characteristics of our study population, with cognitive impairment being severe in all three groups.
223 The NPI group with no significant NPS had the largest proportion of bed-ridden residents. An
224 interesting finding was that the residents with the highest NPS burden at baseline also had the best
225 mobility and the highest number of falls. This seems logical as these residents are physically more
226 active during the day, thus having more opportunities to fall. This is in line with earlier research on
227 fall risk in long-term care.²⁰ However, even after adjustment for mobility, the severity of NPS
228 remained significantly associated with higher incidence of falls, indicating that NPS burden is an
229 independent predictor for falls.

230 The use of psychotropic medication in our study population was very high in all groups. However,
231 there were significant differences in the number of psychotropics used in each NPS group. The
232 group with highest NPS burden also had the highest number of psychotropics (2.3), compared with
233 the groups with low NPS burden (2.1) and no significant NPS (1.8). Several earlier studies suggest
234 that psychotropic medication use increases the risk of falls.³⁶⁻³⁸ To gain more insight into this
235 relationship, we looked at the differences in fall rates among residents with and without
236 psychotropic drug use. In our study, only NPI level was associated with incidence rate of falls,
237 psychotropic drug use was not. There was no interaction indicating that psychotropic drug use did
238 not modify this relationship. Our study suggests that in this special long-term population with
239 severe cognitive impairment, the NPS burden is more important in determining falls than
240 psychotropic drug use.

241 In our study one-third of all falls during the one-year follow-up led to fall related negative
242 consequences (20% to injuries, 7% to hospitalizations, 3% to fractures). This result is consistent
243 with previous studies that have found that most falls do not result in injury.^{4, 8, 20} Even though all
244 falls do not lead to injury, every fall is significant as a previous fall is an important risk factor for
245 another fall.³⁹

246 Our study has some limitations that should be considered when interpreting the results. First, we did
247 not ask about the use of physical restraints. It is well known that, despite clear evidence for a lack of
248 effectiveness and safety, physical restraints are frequently used in nursing homes and their use is
249 associated with falls.⁴⁰⁻⁴¹ Another limitation is that only regularly used psychotropic medication was
250 considered in our study. Psychotropics administered "pro re nata" may have had a different impact
251 on falls and their consequences. Finally, as a longitudinal follow-up study of a special cohort, we
252 cannot rule out unknown confounders having an effect on falls. **Additionally we do not have data on**
253 **past fall history.**

254 Our study has several strengths. The study sample is large and representative of older long-term
255 care residents with cognitive impairment. We used many well-validated assessment instruments,
256 and data were collected by trained study nurses resulting in high data validity. Another important
257 strength is that, to our knowledge, no other study has previously examined the impact of severity of
258 NPS on fall rate, nor has the association between NPS subgroups and fall rate or the interaction of
259 severity of NPS and psychotropics with falls been investigated.

260 **Conclusions and Implications**

261 Most falls are not the result of a single cause, but occur due to an interaction of several risk factors.
262 Thus, a multifactorial approach to fall prevention is recommended. The findings of this study
263 indicate that evaluation of NPS, and especially severity of NPS, and NPS subgroups should be part
264 of the comprehensive assessment when aiming to prevent falls in long-term care residents with
265 cognitive impairment.

266

267 The authors declare that they have no conflicts of interest relevant to this report.

268

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392 **Table 1.** Characteristics of residents grouped by severity of neuropsychiatric symptoms according
393 to Neuropsychiatric Inventory (NPI) total score

394 **Table 2.** Falls and fall-related negative consequences during the 12-month follow-up

395 **Table 3.** Multivariate poisson regression analysis for incidence of falls.

396 **Figure 1.** Incidence of falls per person-years (pyrs) according to Neuropsychiatric Inventory (NPI)
397 total score. Adjusted for age, sex and mobility.

398 **Figure 2.** Association between neuropsychiatric symptoms subgroups and incidence rate ratio
399 (IRR) of falls and fall related negative consequences per 1-SD. Adjusted for age, sex, and mobility.

400 **Figure 3.** Incidence rate of falls during the 12-month follow-up per person-years (pyrs) among
401 residents with and without psychotropic medication according to Neuropsychiatric Inventory (NPI)
402 level.