The relationship between pain and disruptive behaviors in nursing home resident with dementia

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Abstract

Background: Nursing home residents with dementia gradually lose the ability to process information so that they are less likely to express pain in typical ways. These residents may express pain through disruptive behaviors because they cannot appropriately verbalize their pain experience. The objective of this study was to investigate the effect of pain on disruptive behaviors in nursing home residents with dementia.

Methods: This is a secondary analysis of the Minimum Data Set (MDS 2.0) assessment data on long-term care from the state of Florida. The data used in this study were the first comprehensive assessment data from NH residents with dementia aged 65 and older (N = 56,577) in Medicare- or Medicaid-certified nursing homes between January 1, 2009 and December 31, 2009. Variables examined were pain, wandering, aggression, agitation, cognitive impairment, activities of daily living impairments, and demographic characteristics. Ordinal logistic regression was used to evaluate the effect of pain on disruptive behaviors.

Results: Residents with more severe pain are less likely to display wandering behaviors (OR = .77, 95% CI for OR = [0.73, 0.81]), but more likely to display aggressive and agitated behaviors (OR = 1.04, 95% CI for OR = [1.01, 1.08]; OR = 1.17, 95% CI for OR = [1.13, 1.20]).

Conclusions: The relationship between pain and disruptive behaviors depends on the type of behaviors. Pain is positively correlated with disruptive behaviors that do not involve locomotion (e.g., aggression and agitation), but negatively related to disruptive behaviors that are accompanied by locomotion (e.g., wandering). These findings indicate that effective pain management may help to reduce aggression and agitation, and to promote mobility in persons with dementia.

Keywords: Disruptive behaviors, Pain, Dementia, Nursing home

Background

Pain assessment in nursing home (NH) residents with dementia is challenging due to cognitive and communicative impairments. Pain self-report, the gold standard assessment in cognitively intact persons, is questionable in cognitively impaired NH residents because dementia impairs their ability to remember, interpret, and respond to pain [1,2]. NH residents with dementia gradually lose the ability to process information so that they are less likely to express pain in typical ways, even when there is a probable cause for pain [1]. Therefore, pain is often under-reported in NH residents with dementia. These residents may express pain through disruptive behaviors [3], because they cannot appropriately verbalize their pain experience.

Disruptive behaviors, also known as “problematic behaviors,” “disturbing behaviors,” or “challenging behaviors,” refer to inappropriate, repetitive, or dangerous behaviors that are disruptive to the living and working environment in the NH [4,5]. Among many disruptive behaviors, three behaviors are most prominent in the current literature: wandering behaviors, aggressive behaviors, and agitated behaviors [6,7]. Wandering occurs in approximately 40 to 60% of NH residents with dementia [8], and aggression and agitation occurs in about 50% to 80% of NH residents with cognitive impairments [9].
Disruptive behaviors are problematic to NH residents and staff. Disruptive behaviors are associated with injuries and hospitalizations among NH residents with dementia, and contribute to stress and burnout among caregivers [10,11]. The cost of care for NH residents with dementia is three times higher than that of other NH residents, and about 30% of these costs are attributed to the management of disruptive behaviors [12]. Psychoactive medications or restraints are often used to manage disruptive behaviors [13]; however, these often lead to falls, impaired functioning, and decreased mobility. The use of restraints is also an affront to personal dignity. The better approach to managing disruptive behaviors is to control their possible causes, such as pain.

Thus, the purpose of this study is to explore the relationship between pain and disruptive behaviors in NH residents with dementia. Such information may identify potential new intervention approaches for managing these behaviors.

### Theoretical framework

The Need-driven Dementia-compromised Behavior (NDB) model [14] was used to guide this study of the relationship between pain and disruptive behaviors in NH residents with dementia (Figure 1). The NDB model posits two main constructs that are associated with dementia-compromised behaviors: background factors and proximal factors. Background factors represent those characteristics that place older adults at risk for disruptive behaviors. Proximal factors represent the conditions under which disruptive behaviors occur. We conceptualized pain as a proximal factor that would have a direct relationship with disruptive behaviors (e.g., wandering, aggression, and agitation). For this study, the level of cognitive impairment, activities of daily living (ADL) impairment, and demographic characteristics (e.g., age and sex) represent background factors. These variables were selected as covariates because they have established relationships with both pain and disruptive behaviors [15-18] and may influence the relationship between pain and disruptive behaviors.

### Methods

This is a secondary analysis of the nursing home Minimum Data Set (MDS) from the state of Florida during calendar year 2009. The first comprehensive assessment for each NH resident was used in this study. The archived data files of the most recent version of MDS (MDS 3.0) were not yet available to researchers, but are due for release in early 2013 [19]. The MDS data are mandatory in all NHs certified to participate in Medicare and Medicaid. Approval for the study was obtained from the University of Florida Health Science Center Institutional Review Board.

The MDS assessment data, standardized data on residents’ status based on routine and continuous observations by nursing staff, provides comprehensive information on all the NH residents. The MDS assessment is completed on admission to the facility, on a quarterly basis thereafter, and upon significant changes in status [20]. The complete federal database consists of over 1.5 million older adults who live in NHs throughout the United States. Although it is used primarily for clinical purposes, the MDS has also been used for research on cognition and behavioral symptoms in this population [21-23]. Several MDS subscales have been created and evaluated, and have demonstrated acceptable reliability and validity: MDS-Pain severity scale [24], MDS-Depression Rating Scale [25], MDS-Aggression Behavior Scale [26], MDS-Challenging Behavior Profile [27], MDS-Discomfort Behavior Scale [28], MDS-Cognitive Performance Scale [28,29], MDS-index of social engagement [30,31], MDS-Activities of Daily Living scale [32,33], Resident Assessment Instrument-Mental Health [34], and MDS-Change in Health, End-stage disease and Signs and Symptoms [35]. Details of the reliability and validity coefficients for each of the major study variables are described in the measurement section.

### Data used in this study

The data used in this study were collected on residents with dementia in Medicare- or Medicaid-certified NHs who have a MDS comprehensive assessment on file. The data were acquired from the Centers for Medicare & Medicaid Services. Selection criteria were applied to ascertain data.
from NH residents older than 65 years old with Alzheimer’s disease or other dementia, based on documented medical diagnosis. Data from comatose residents were excluded, because these residents cannot display the disruptive behaviors investigated in this study. This selection process yielded 56,577 unique cases for the analyses.

The sample is mostly female (67.7%), and a mean age of 84 (years range = 65–109). The prevalence of disruptive behaviors is as follows: wandering behaviors (9.0%), aggressive behaviors (24.4%), and agitated behaviors (24.1%) (Table 1).

**Measurement**

MDS subscales and items were used to indicate the main study concepts: pain, wandering, aggression, and agitation. These are described below.

**Pain**

The MDS-pain severity scale [24], combining both pain frequency (0 = no pain, 1 = pain less than daily, and 2 = pain daily) and pain intensity (1 = mild pain, 2 = moderate pain, and 3 = horrible or excruciating pain), was used to assess pain severity in NH residents with dementia. This scale can range from 0 to 3, with higher scores indicating greater pain severity. NH residents’ self-report is reflected in the MDS pain items if residents can self-report and staff completing the MDS assessments have confidence in residents’ self-report. Otherwise, the staff who complete the MDS assessment document pain symptoms based on proxy reports from facility nursing staff that provides care to the residents. The MDS-pain severity scale has been reported to have an inter-rater reliability coefficient of 0.73, and kappa coefficient of 0.70 with a Visual Analogue Scale in a study involving 95 U.S. nursing home residents at 25 Medicare-certified skilled nursing facilities in Massachusetts [24].

**Disruptive behaviors**

The MDS-wandering item was used to measure the frequency of wandering in the last 7 days. Wandering frequency is recorded by staff observation. It is recorded as no wandering, wandering occurred 1 to 3 days, wandering occurred 4 to 6 days, and daily wandering. The wandering item has been reported to have a reliability coefficient of 0.63, and an inter-rater reliability of 0.95 [36,37].

The MDS-Aggression Behavior Scale (MDS-ABS) was used to measure the frequency of aggressive behaviors. The MDS-ABS is a sum score of four MDS items: verbally abusive behavioral symptoms, physically abusive behavioral symptoms, socially inappropriate behavioral symptom, and resisting care. The MDS-ABS can range from 0 to 12, with higher scores indicating more frequent aggressive behaviors. The MDS-ABS has been reported to have an internal consistency reliability of 0.79 to 0.95, and a criterion validity coefficient of 0.72 with Cohen-Mansfield Agitation Inventory aggression subscale scores [26].

The revised MDS-Challenging Behavior Profile (MDS-CBP) agitation subscale was used to assess the frequency of agitated behaviors. The revised agitation scores, calculated using two MDS items (e.g., periods of restlessness and repetitive physical movements), can range from 0 to 3, with higher scores indicating more frequent agitated behaviors. This revised agitation scale has Cronbach’s alpha coefficient of .68. The original MDS-CBP agitation subscale, computing from 4 MDS items (e.g., periods of restlessness, repetitive physical movements, wandering, and socially inappropriate behavioral symptom), has been reported to have Cronbach’s alpha coefficient of 0.70, inter-rater reliability of 0.61, and a Spearman’s rank correlation coefficient of 0.50 with Behavior Rating Scale for Psychogeriatric Inpatients [27].

### Table 1 Sample characteristics

| Characteristic          | Number | Total sample |
|-------------------------|--------|--------------|
| Age, mean ± SD          | 56577  | 84.37 ± 7.43 |
| Gender, n (%)           |        |              |
| Male                    | 18,265 | (32.3)       |
| Female                  | 38,301 | (67.7)       |
| MDS-CPS, mean ± SD      | 56543  | 3.17 ± 1.52  |
| MDS-ADL, mean ± SD      | 56577  | 18.66 ± 6.41 |
| Pain severity, mean ± SD| 56568  | 0.48 ± 0.70  |
| Wandering behaviors, n (%)| 56573 |              |
| No wandering (MDS-wandering = 0) | 51,463 | (91.0) |
| 1-3 days in 7 days (MDS-wandering = 1) | 2,637 | (4.7) |
| 4-6 days in 7 days (MDS-wandering = 2) | 994 | (1.8) |
| Wandering daily (MDS-wandering = 3) | 1,479 | (2.6) |
| Aggressive behaviors, n (%)| 56572 |              |
| None (MDS-ABS = 0)      | 42,764 | (75.6)       |
| Moderate (MDS-ABS = 1 – 2) | 9,667 | (17.1)       |
| Severe (MDS-ABS = 3 – 5) | 3,390 | (6.0)        |
| Very severe (MDS-ABS = 6 – 12) | 751 | (1.3)        |
| Agitated behaviors, n (%)| 56571 |              |
| None (revised MDS-CBP agitation = 0) | 42,941 | (75.9) |
| Mild (revised MDS-CBP agitation = 1) | 6,916 | (12.2)       |
| Moderate (revised MDS-CBP agitation = 2) | 5,099 | (9.0)        |
| Severe (revised MDS-CBP agitation = 3) | 1,615 | (2.9)        |

MDS-CPS = MDS-Cognitive Performance Scale.
MDS-ADL = MDS-Activities of Daily Living impairment scale.
MDS-ABS = MDS-Aggression Behavior Scale.
Revised MDS-CBP agitation = revised MDS-Challenging Behavior Profile, agitation subscale.
Background factors

The MDS-cognitive performance scale (MDS-CPS) [38] was used to measure the level of cognitive impairment. The MDS-CPS score is calculated using five MDS items: comatose, short-term memory, cognitive skills or daily decision making, making oneself understood, and self-performance in eating. The MDS-CPS can range from 0 to 6, with higher scores indicating more cognitive impairment. The MDS-CPS has been reported to have a kappa coefficient of 0.45-0.75 with Mini-Mental State Examination, a kappa coefficient of 0.41-0.77 against Global Deterioration Scale, a kappa coefficient of 0.66 against Psychogeriatric Dependency Rating Scale, a kappa coefficient of 0.45 against Mattis Dementia Rating Scale [29,38-41].

The MDS-Activities of Daily Living-Long Form (MDS ADL-Long Form) [42] was used to measure the level of ADL impairment. The MDS ADL-Long Form scores are calculated using 7 MDS items: self-performance of bed mobility, transfer, locomotion on unit, dressing, eating, toilet use, and personal hygiene. MDS ADL-Long Form can range from 0 to 28, with higher scores indicating more impairment of ADLs. The MDS ADL-Long Form has been reported to have a reliability coefficient of 0.92-0.97, an inter-rater reliability coefficient of 0.61-0.95, and a kappa coefficient of 0.58 – 0.79 against Physical Self-Maintenance Scale [31,43].

Demographics characteristics (e.g., age and gender) were collected from the MDS form. Age was a continuous variable and gender was dichotomous (0 = female; 1 = male). They were included as covariates in the analyses.

Statistical analysis

Analyses were performed using SPSS, version 20 (IBM Inc., Armonk, NY). Multivariate analyses were conducted to explore the relationship between pain and disruptive behaviors in this sample. Aggression was severely positively skewed, and none of the transformations (e.g., logarithmic transformation, square root transformation, inverse transformation, and square transformation) resolved the normal distribution issue. Therefore, aggression was collapsed into four groups (none, moderate, severe, and very severe), based on published algorithms in the literature [26]. Aggression was transformed as none (MDS-ABS = 0), moderate (MDS-ABS = 1–2), severe (MDS-ABS = 3–5), and very severe (MDS-ABS = 6–12). Due to concerns that NH residents who take psychotropic medications (e.g., antipsychotics, antidepressants, etc.) may exhibit less frequent disruptive behaviors [44], we re-ran the statistical analysis excluding these subjects.

Since the level of measurement of the dependent variables was ordinal, logistic regression for ordinal variables was used to evaluate the effect of pain severity on the three disruptive behaviors, after controlling for covariates. Using the same independent variables in analysis with different dependent variables carries the risk of inflating the Type I error. To keep the overall risk of a Type I error to the 5% level, p-value for each regression analysis is set at .017.

Results

The results of ordinal logistic regression on three disruptive behaviors, after controlling for covariates (e.g., the level of cognitive impairment, the level of ADL impairment, and sociodemographic factors) are described below.

| Variables | Wandering | | | Aggression | | | Agitation | | |
|---|---|---|---|---|---|---|---|---|---|
| Independent Variable | B | OR | 95% CI for OR | B | OR | 95% CI for OR | B | OR | 95% CI for OR |
| Pain | −0.26* | 0.77 | [0.73, 0.81] | 0.04* | 1.04 | [1.01, 1.08] | 0.15* | 1.17 | [1.13, 1.20] |
| Covariates | | | | | | | | | |
| MDS-CPS | 0.68* | 1.97 | [1.91, 2.02] | 0.36* | 1.43 | [1.41, 1.46] | 0.46* | 1.58 | [1.55, 1.60] |
| MDS-ADL | −0.15* | 0.87 | [0.86, 0.87] | −0.03* | 0.98 | [0.97, 0.98] | −0.02* | 0.98 | [0.97, 0.98] |
| Age | −0.01* | 0.99 | [0.98, 0.99] | −0.01* | 0.99 | [0.99, 0.99] | −0.01* | 0.99 | [0.99, 0.99] |
| Sex | | | | | | | | | |
| Male | 0.22* | 1.25 | [1.17, 1.33] | 0.28* | 1.33 | [1.27, 1.39] | 0.24* | 1.27 | [1.22, 1.33] |
| Female | 0.00 | 1.00 | 0.00 | 1.00 | 1.00 | | | |

Nagelkerke R-square: Wandering = 0.15, Aggression = 0.06, Agitation = 0.08. B = logistic regression coefficient, OR = Odds Ratio = \( \text{Exp}(B) \), MDS-CPS = MDS-Cognitive Performance Scale. MDS-ADL = MDS-Activities of Daily Living impairment scale. *p < .001.
The effect of pain on aggressive behaviors
Pain severity is positively associated with the frequency of aggressive behaviors (Table 2). NH residents with more severe pain are more likely to display aggressive behaviors (Logistic regression coefficient = 0.04, p < .001, Odds Ratio = 1.04, 95% CI for Odds Ratio = [1.01, 1.08]).

The effect of pain on agitated behaviors
Pain severity is positively associated with the frequency of agitated behaviors (Table 2). NH residents with more severe pain are more likely to display agitated behaviors (Logistic regression coefficient = 0.15, p < .001, Odds Ratio = 1.17, 95% CI for Odds Ratio = [1.13, 1.20]).

The study results in subsample without psychotropic medications
The results of ordinal logistic regression in the subsample without psychotropic medications (e.g., antipsychotics, antidepressants, etc.) are summarized in Table 3. These results of ordinal logistic regression are similar when NH residents who used psychotropic medications in the past 7 days were excluded. Pain severity is negatively associated with the frequency of wandering behaviors, but positively associated with the frequency of aggressive and agitated behaviors.

Discussion
It was found that more severe pain is associated with less frequent wandering behaviors, but more frequent aggressive and agitated behaviors, after controlling for covariates. Most of the published literature suggested that there is a positive relationship between pain and disruptive behaviors in general [6,11,45]. However, the results of this study suggest that the relationship between pain and disruptive behaviors depends on the type of behaviors examined. The direction of the relationship between these variables depends on whether the disruptive behaviors are accompanied by locomotion.

Pain is positively correlated with disruptive behaviors that do not involve locomotion (e.g., aggression and agitation), but negatively related to disruptive behaviors that are accompanied by locomotion (e.g., wandering). That is, residents who experience more severe pain are more likely to display aggression and agitation, and less likely to move around.

The finding that pain and aggressive or agitated behaviors are positively linked in NH residents with dementia is consistent with other published reports. Buffum and colleagues [46] reported that pain was positively related to agitation \( (r = .50, p = .003) \) using a bivariate correlation analysis in 33 Veterans Affairs NH residents with dementia. Manfredi and colleagues [47] demonstrated that opioid treatment for pain reduced agitation in 13 NH residents with dementia who were more than 85 years old (mean change in CMAI score: -6.4, 95% CI [-10.96, -1.8]). Both of these studies have a small sample size. Thus, the results of this study using a large sample from all the nursing home residents with dementia in the state of Florida substantiates and extends the positive relationship between pain and non-locomotive disruptive behaviors from these previous findings.

In contrast, the finding on the relationship between pain and wandering behavior in this study is opposite to the findings presented in the literature review. Kiely and colleagues [48] used MDS assessment data from 8,982 NH residents, and reported that NH residents who expressed sadness or pain in MDS assessment data were 65% more likely to develop wandering behaviors than their counterparts who did not express sadness or pain (OR = 1.65, p = .02). Our study measured pain more specifically using the MDS-pain severity scale [24], combining both pain frequency and pain intensity, while Kiely and the colleagues [48] measured pain by a dichotomized expression of sadness or pain. Sadness is not typically considered an indicator of

| Variables | Wandering |  |  | Aggression |  |  | Agitation |  |  |
|-----------|-----------|---|---|-----------|---|---|-----------|---|---|
|           | B OR 95% CI for OR | B OR 95% CI for OR | B OR 95% CI for OR |
| Pain      | −0.33 0.72 [0.63, 0.83] | 0.07* 1.07 [1.01, 1.15] | 0.15* 1.16 [1.08, 1.25] |
| Covariates |  |  |  |
| MDS-CPS   | 0.65 1.87 [1.76, 2.00] | 0.29* 1.34 [1.29, 1.38] | 0.42* 1.53 [1.47, 1.58] |
| MDS-ADL   | −0.15 0.86 [0.85, 0.87] | −0.04 0.97 [0.96, 0.97] | −0.04 0.96 [0.96, 0.97] |
| Age       | 0.00 1.00 [0.99, 1.01] | 0.01* 1.01 [1.00, 1.02] | 0.00 1.00 [1.00, 1.01] |
| Sex       |  |  |  |
| Male      | 0.12 1.12 [0.96, 1.31] | 0.19* 1.21 [1.10, 1.33] | 0.22* 1.24 [1.13, 1.37] |
| Female    | 0.00 1.00 [0.99, 1.01] | 0.00 1.00 [1.00, 1.01] | 1.00 |

Nagelkerke R-square: Wandering = 0.13, Aggression = 0.03, Agitation = 0.05. B = logistic regression coefficient, OR = Odds Ratio = Exp(B). MDS-CPS = MDS-Cognitive Performance Scale. MDS-ADL = MDS-Activities of Daily Living impairment scale. *p < .001. **p < .05.
pain, and its inclusion may have confounded pain and depression or mood disorder.

Several limitations of this study should be noted. First, this study is inherently limited by secondary analysis of federally mandated MDS assessment data, and the effect of clustering within facility is not controlled in this study. The variables and the procedures cannot be controlled. The MDS assessment data may have some variability due to different styles and skills of MDS coordinators in each facility. Second, the role of pain medications is not considered in this study. The highest level of pain could have been managed by pain medications, but it is not possible to discern this in the MDS assessment data. However, similar to our study, most of the literature reported the relationship between highest level of pain and the frequency of behavioral symptoms during the observation period without controlling for pain medications [11,49]. Third, the amount of variance in disruptive behaviors that is explained by these logistic regression models is small (ranging from 6% to 15%). This suggests that there are other factors that contribute to disruptive behaviors that were not specified in our models. Finally, this study design is descriptive and cross-sectional. As such, this study is not able to examine causal relationships between pain and disruptive behaviors.

Findings from this study can be a foundation for future research. Studies using prospective designs are needed to validate these findings. Also, randomized controlled trials can be used to compare comprehensive pain management and usual pain management with regard to the frequency of disruptive behaviors. This type of study can provide evidence for causal relationships between pain management and disruptive behaviors and support changes in clinical practice. Third, future research would include the longitudinal MDS assessment data to examine trends over time. The longitudinal nature of MDS assessment data, collected every three months or more often, provides an opportunity to describe change over time, and facilitates the use of more powerful statistical analysis techniques to describe both within- and between-person changes.

Conclusions

Pain exacerbated disruptive behaviors that are not locomotion-based. In order to reduce these disruptive behaviors, their underlying causes, such as pain, should be investigated and well managed. However, pain assessment in cognitively impaired residents can be challenging. Comprehensive pain assessment should be developed further, and pain should be well controlled to reduce these problematic disruptive behaviors.

Competing interests

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Authors’ contributions

HA conceptualized the study, completed all statistical analyses, and wrote the manuscript. AH provided oversight and consultation during all aspects of the study. Both authors read and approved the final manuscript.

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