This form contains the measure information submitted by stewards. Blank fields indicate no information was provided. Attachments also may have been submitted and are provided to reviewers. The subcriteria and most of the footnotes from the evaluation criteria are provided in Word comments within the form and will appear if your cursor is over the highlighted area. Hyperlinks to the evaluation criteria and ratings are provided in each section.

**TAP/Workgroup** (if utilized): Complete all highlighted areas of the form. Evaluate the extent to which each subcriterion is met. Based on your evaluation, summarize the strengths and weaknesses in each section.

**Note:** If there is no TAP or workgroup, the SC also evaluates the subcriteria (highlighted areas).

**Steering Committee:** Complete all highlighted areas of the form. Review the workgroup/TAP assessment of the subcriteria, noting any areas of disagreement; then evaluate the extent to which each major criterion is met; and finally, indicate your recommendation for the endorsement. Provide the rationale for your ratings.

---

**MEASURE DESCRIPTIVE INFORMATION**

<table>
<thead>
<tr>
<th>Measure Title</th>
<th>Percent of Residents with Pressure Ulcers That Are New or Worsened (Short-Stay)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brief description of measure</td>
<td>This measure updates CMS' current QM pressure ulcer measure which currently includes Stage 1 ulcers. The measure is based on data from the MDS 3.0 assessment of short-stay nursing facility residents and reports the percentage of residents who have Stage 2-4 pressure ulcers that are new or have worsened. The measure is calculated by comparing the Stage 2-4 pressure ulcer items on the discharge assessment and the previous MDS assessment (which may be an OBRA admission or 5-day PPS assessment). The quality measure is restricted to the short-stay population defined as those who are discharged within 100 days of admission. The quality measure does not include the long-stay residents who have been in the nursing facility for longer than 100 days. A separate measure has been submitted for them.</td>
</tr>
</tbody>
</table>

**Type of Measure:** Outcome

---

**CONDITIONS FOR CONSIDERATION BY NQF**

Four conditions must be met before proposed measures may be considered and evaluated for suitability as voluntary consensus standards:

A. The measure is in the public domain or an intellectual property (measure steward agreement) is signed. **Public domain only applies to governmental organizations. All non-government organizations must sign a measure steward agreement even if measures are made publicly and freely available.**

A.1 Do you attest that the measure steward holds intellectual property rights to the measure and the right to use aspects of the measure owned by another entity (e.g., risk model, code set)? **Yes**
### A.2 Indicate if Proprietary Measure (as defined in measure steward agreement):

| B | Y | N |

### A.3 Measure Steward Agreement: Government entity and in the public domain - no agreement necessary

### A.4 Measure Steward Agreement attached:

| B | Y | N |

### B. The measure owner/steward verifies there is an identified responsible entity and process to maintain and update the measure on a schedule that is commensurate with the rate of clinical innovation, but at least every 3 years. Yes, information provided in contact section

### C. The intended use of the measure includes both public reporting and quality improvement.

**Purpose:** Public reporting, Internal quality improvement

### D. The requested measure submission information is complete. Generally, measures should be fully developed and tested so that all the evaluation criteria have been addressed and information needed to evaluate the measure is provided. Measures that have not been tested are only potentially eligible for a time-limited endorsement and in that case, measure owners must verify that testing will be completed within 12 months of endorsement.

**D.1 Testing:** No, testing will be completed within 24 months

**D.2 Have NQF-endorsed measures been reviewed to identify if there are similar or related measures?** Yes

### 1. IMPORTANCE TO MEASURE AND REPORT

**1a. High Impact**

**Ev Rat ing**

| C | P | M | N |

**1a.1 Demonstrated High Impact Aspect of Healthcare:** Severity of illness, Patient/societal consequences of poor quality, High resource use

**1a.2**

**1a.3 Summary of Evidence of High Impact:** Pressure ulcers are serious medical conditions. They typically result from prolonged periods of uninterrupted pressure on the skin, soft tissue, muscle, and bone. Patients are often elderly, stroke and diabetic patients; those with dementia, circulatory diseases, dehydration, and malnutrition; and people who use wheelchairs or are bedridden—those is, any patient with impaired mobility or sensation. Pressure ulcers interfere with the activities of daily living, predispose patients to osteomyelitis and septicemia, and are strongly associated with longer hospital stays and mortality.

Pressure ulcers are high-volume and high-cost adverse events across the spectrum of health care settings from acute hospitals to home health. The prevalence of pressure ulcers in health care facilities is increasing, with some 2.5 million patients being treated annually for pressure ulcers in acute care facilities. In 2006, there were 503,300 acute hospital stays during which pressure ulcers were noted—a 78.9% increase.
from 1993 when there were about 281,300 hospital stays related to pressure ulcers. (1, 5)

Pressure ulcer incidence rates vary considerably by clinical setting—ranging from 0.4% to 38% in acute care, from 2.2% to 23.9% in skilled nursing facilities and nursing homes, and from 0% to 17% in home care. (3, 16)

Patients with acute care hospitalizations related to pressure ulcers were more likely to be discharged to long-term care facilities (e.g., a skilled nursing facility, an intermediate care facility, or a nursing home), than hospitalizations for all other conditions. (3, 4) In fact, more than half of principal pressure ulcer stays (53.4%) and secondary pressure ulcer stays (54.5%) were discharged to long-term care—more than 3 times the rate of hospitalizations for all other conditions (16.2%). (4)

Pressure ulcers are one of the most important measures of the quality of clinical care in nursing facilities. The CDC conducts the National Nursing Home Survey, a continuing series of national sample surveys of nursing homes, their residents, and their staff. Data for the survey were obtained through personal interviews with facility administrators and designated staff who used administrative records to answer questions about the facilities, staff, services and programs, and medical records to answer questions about the residents. A total of 1,174 nursing home facilities participated in the latest National Nursing Home Survey. (6)

As reported in the 2004 National Nursing Home Survey results, about 159,000 current U.S. nursing home residents (11%) had pressure ulcers. Stage 2 ulcers were the most common, accounting for about 50% of all pressure ulcers. Stages 1, 3, and 4 made up about the other 50% of all pressure ulcers. (6)

Stage 1 pressure ulcers are not included in the proposed quality measure, researchers have suggested that inclusion of Stage 1 pressure ulcers in the quality measures adds little value. (17)

Graph is shown in the attached Figure 1: Percentage of Nursing Home Residents with Pressure Ulcers: United States, 2004.

In 2006, Abt Associates examined pressure ulcer incidence and prevalence across post-acute settings. For nursing homes, MDS 2.0 assessments were used for April 1, 2006, through July 15, 2006. The prevalence of pressure ulcers Stage 1-4 was 13%, with the prevalence of Stage 3-4 ulcers being 3% nationwide. (4)

Pressure ulcers may cause extreme discomfort to the patient and often lead to serious, life-threatening infections, which substantially increase the total cost of care. (1, 7, 8) The main driver of cost is the presence of complications, which involve diagnostic tests, additional monitoring, more expensive pressure-relieving surfaces, and extended length of stays. (7)

As reported in the Federal Register, in 2006 there were 322,946 reported cases of Medicare patients with a pressure ulcer as a secondary diagnosis—each case had an average charge of $40,381 for a hospital stay, for an annual total cost of $13 billion. (9) The Advancing Excellence in America’s Nursing Homes Campaign, a national effort launched in 2006 to help nursing homes measurably to improve care, reported that it can cost as much as $19,000 to treat a single Stage 4 pressure ulcer. (10)

To address this critical clinical issue, there are numerous national healthcare organizations with ongoing efforts and publications to prevent pressure ulcers, monitor prevalence, and improve treatment. Examples listed below are representative but not exhaustive and include the following:

• The Joint Commission on Accreditation of Healthcare Organizations offers National Patient Safety Goals for Long-Term Care. (11)
• The Institute for Healthcare Improvement’s 5 Million Lives Campaign was a voluntary initiative to protect patients from 5 million incidents of medical harm over the 2006-2008 period. (12)
• CDC’s National Center for Health Statistics and National Nursing Home Survey monitor pressure ulcer prevalence. (6)
• The On-Time Quality Improvement for Long-Term Care Program, funded by the Agency for Healthcare Research and Quality (AHRQ) in collaboration with State departments of health, is a Quality Improvement Organization (QIO), or a trade association, to improve nursing home care. This national effort focuses on prevention and timely treatment of pressure ulcers during routine care. New tools to document pressure ulcer healing and treatments and reports to help monitor the healing process have been developed as part of the On-Time Quality Improvement Program. (4, 13)
• The National Pressure Ulcer Advisory Panel (NPUAP) uses research, public policy, and education to improve patient outcomes in pressure ulcer prevention. (14)
• The NQF sponsors the “National Voluntary Consensus Standards for Developing a Framework for Measuring
Quality for Prevention and Management of Pressure Ulcers initiative. This CMS-funded project began in 2008 and has several objectives, including determining how to measure the incidence and prevalence of pressure ulcers and their staging and harmonizing measure specifications across settings of care.(4, 15)

- The Advancing Excellence Campaign is a national campaign to encourage, assist, and empower nursing homes to improve the quality of care and life for residents. Of the eight goals set by the effort, the fourth goal is for nursing home residents receive appropriate care to prevent and appropriately treat pressure ulcers when they develop.(10)


1b. Opportunity for Improvement

1b.1 Benefits (improvements in quality) envisioned by use of this measure: This measure is intended to encourage nursing facilities to focus on this important clinical issue in order to prevent pressure ulcers and to closely monitor and promote healing of existing pressure ulcers.

1b.2 Summary of data demonstrating performance gap (variation or overall poor performance) across providers:
The short-stay pressure ulcer measure is one of the current CMS publicly reported quality measures for nursing facilities and Five-Star Quality Rating System. In its analysis of the current quality measure using MDS 2.0 data from 2006, the University of Colorado found variability across facilities in the rates of pressure ulcers for short-stay residents, suggesting that it is possible for facilities to improve. As presented in the table below, the national overall nursing facility mean was 17.1% and the standard deviation was 9.3%. The short-stay pressure ulcer quality measure demonstrated significant variability across facilities; from 7.0% at the 10th percentile to 28.7% at the 90th percentile with only 1.1% of facilities reporting short-stay residents with no pressure ulcers.

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<tr>
<th>1</th>
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<td>B</td>
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See attached Table 1: Measure Variability Across Facilities.
The most recent state and national averages for the current MDS 2.0 pressure ulcer quality measure are reported on CMS's Nursing Home Compare Web site for the target quarter ending in June 2009. The data continue to demonstrate the ongoing gap in facility performance; the national average was 16.9%, and state averages ranged from a low of 11.3% to a high of 25.6%. The Advancing Excellence Campaign in America’s Nursing Homes is a national effort begun in 2006 to encourage, assist, and empower nursing facilities to improve the quality of care and life for residents. The coalition comprises long-term care providers, medical professionals, consumers, employees, and state and federal agencies and is the largest and first coalition of its kind to measure quality by setting clinical and organizational goals for nursing facilities. As of October, 2009, the Advancing Excellence Campaign has recruited over 7,600 nursing facilities—47% of all nursing homes in the United States. Of the eight goals set by the effort, the fourth goal is for nursing facility residents to receive appropriate care to prevent and appropriately treat pressure ulcers when they occur. Although the current focus of the campaign is on high-risk, long-stay residents, the goals and results support the evidence of high impact and clinical significance.

As stated in the Implementation Guide, for this goal, the following objectives have been set for December 31, 2011 as part of Phase 2 efforts:

A: The national average for pressure ulcers will be at or below 9%.
B: 30% of nursing facilities will report rates of pressure ulcers at or below 6%.
C: The average of the scores of the nursing facilities exceeding the 2009 Q1 90th percentile (n = 1147) will be reduced from 25% to 18%.
D: Compared with June 2009, there will be 3,000 fewer residents with pressure ulcers per 100,000 nursing facility residents. Applying this to the current pressure ulcer denominator of approximately 750,000 results in 225,000 fewer residents with pressure ulcers.
E: Each state will attain an average facility-level improvement of 1 decile.
F: Nursing facilities will set a specific target to improve the prevalence of pressure ulcers by 1 decile rank over the next 24-month period.

To date, progress has been steady but incremental in meeting these goals as demonstrated by campaign objective graphs: http://www.nhqualitycampaign.org/files/reports/results/q2-2009/Goal1_NationalObjectives_2009Q2_1page.pdf (4). As previously stated, although the initial focus of this campaign is long-stay residents, the improvements in the nursing facilities involved in this campaign should benefit the all residents in the facility and demonstrates the capability of facilities to improve on this quality.
1b.3 Citations for data on performance gap:

1b.4 Summary of Data on disparities by population group:
Research suggests racial disparities in quality of care in nursing facilities between African Americans and Caucasians (1, 2, 3, 4, 5) and between Hispanics and Caucasians.(6) In 1999, Lapane and colleagues found African American residents, compared with Caucasian residents, had a lower prevalence of early-stage pressure ulcers but a higher prevalence of later stage pressure ulcers (even when controlling for other patient sociodemographic and clinical variables).(5) However, in 2009, CDC reported in their key findings from the 2004 National Nursing Home Survey that there was no significant difference between white and nonwhite populations with respect to having pressure ulcers.(7) No research has been conducted on other types of disparities (e.g., ethnicity, rural/urban, or income) specifically for this measure.

1b.5 Citations for data on Disparities:

1c. Outcome or Evidence to Support Measure Focus

1c.1 Relationship to Outcomes (For non-outcome measures, briefly describe the relationship to desired outcome. For outcomes, describe why it is relevant to the target population): The desired outcome is for a low percentage of short-stay, nursing facility residents to have Stage 2-4 pressure ulcers that are new, worsen, or do not show improvement. Pressure ulcers are a significant clinical concern in the nursing facility population given that those at risk for developing pressure ulcers include the elderly, stroke and diabetic patients, those with dementia, and people who use wheelchairs or who are bedridden—that is, any patients with impaired mobility or sensation.(1, 3, 4) The most recent state and national averages for the current
pressure ulcer quality measure are reported on CMS’s Nursing Home Compare Web site for the target quarter ending in June 2009. Additionally, there is an increase in pressure ulcers in acute care facilities. In 2006, there were 503,300 acute hospital stays during which pressure ulcers were noted. Acute care hospitalizations related to pressure ulcers were more likely to be discharged to long-term care facilities (e.g., skilled nursing facilities, intermediate care facilities, or nursing homes), as compared with hospitalizations for all other conditions. In fact, more than half of principal pressure ulcer stays (53.4%) and secondary pressure ulcer stays (54.5%) were discharged to long-term care—more than 3 times the rate of hospitalizations for all other conditions (16.2%).


1c.2-3. Type of Evidence: Evidence-based guideline, Observational study, Systematic synthesis of research, Cohort study, Expert opinion

1c.4 Summary of Evidence (as described in the criteria; for outcomes, summarize any evidence that healthcare services/care processes influence the outcome):
The evidence identifying risk factors, prevention practices, intervention, and treatment for pressure ulcers is well documented and supported by a substantial number of studies and practice guidelines across health care settings, clinical disciplines, and health care organizations. Risk factors include being elderly, frail, having a diagnosis of stroke, diabetes or dementia, and using a wheelchair for mobility or being bedridden—that is, having impaired mobility, sensation, or circulation, or malnutrition, or weight loss. Pressure ulcers cause considerable harm to patients, hindering functional recovery, often causing pain, and often serving as vehicles for the development of serious infections. They have also been associated with an extended length of stay and increased mortality. In fact, an estimated 60,000 patients die each year from complications resulting from hospital-acquired pressure ulcers.

Pressure ulcers are frequently preventable. Pressure ulcer prevention is not a new concept to health care practitioners. Prevention strategies are recommended in medical, nursing, and physical therapy practice guidelines. The Agency for Healthcare Research and Quality conducted a systematic review of the literature for evidence regarding interventions that could prevent pressure ulcers. The review identified 40 systematic reviews and 45 individual studies that investigated interventions for the prevention of pressure ulcers. Most interventions included topic education, pressure redistribution, and skin care. The effectiveness of the interventions was assessed in 41 studies. The review concluded that interventions were effective in preventing pressure ulcers in most settings. The interventions that were most effective were those that involved education and that included pressure redistribution and skin care. The interventions that were least effective were those that did not include all three components. The review also identified several limitations of the studies, including small sample sizes, lack of blinding, and lack of randomization. The review concluded that interventions to prevent pressure ulcers are effective, but more research is needed to determine the optimal intervention.
care facilities. Many organizations develop prevention guidelines which entail two major steps: identifying patients at risk; and, reliably implementing prevention strategies for all patients who are identified as being at risk.

For years, facilities developed pressure ulcer prevention programs using proven techniques, however, facilities often lack reliable strategies, as well as a long-term commitment, to design caregiver’s work so that prevention remains a priority. (3, 4)

Prevention strategies include six key elements: (1) conducting a pressure ulcer admission assessment for all patients, (2) reassessing risk for all patients daily, (3) inspecting skin daily, (4) managing moisture, (5) optimizing nutrition and hydration, and (6) minimizing pressure. (3, 4, 5, 6)

For existing pressure ulcers, proven treatment strategies are consistent with the Advancing Excellence Campaign in America’s Nursing Homes’ Implementation Guide. (4) The guide provides evidence-based approaches to minimize pressure ulcers. Steps to address this issue include (1) recognition and assessment, (2) cause identification, (3) management, and (4) monitoring. (4) The treatment goal may be either healing, palliative, or maintenance. Pressure ulcer treatment should include wound cleansing and may also include surgical repair, adjunctive therapy, or debridement. (6) Pressure ulcer assessment and treatment should be also documented on admission, on an ongoing basis, and prior to any transition from one health care setting to another. (6)

Education and training for staff on identifying pressure ulcer risk, prevention, and treatment are also critical to facility success and needs to be done routinely to keep staff competent and current. Education must be based on the needs of the staff and appropriate to the patient population. (6)

1c.5 Rating of strength/quality of evidence (also provide narrative description of the rating and by whom): The body of evidence has not been rated.

1c.6 Method for rating evidence:

1c.7 Summary of Controversy/Contradictory Evidence: No contradictory evidence has been identified.


1c.9 Quote the Specific guideline recommendation (including guideline number and/or page number): The clinical guidelines are too extensive to quote here but can be found in citations 1, 2, 3, 4, 5, and 6 below.


1c.11 National Guideline Clearinghouse or other URL: http://www.guideline.gov/browse/browsemode.aspx?node=47462&type=1

1c.12 Rating of strength of recommendation (also provide narrative description of the rating and by whom): The guidelines utilized various rating systems to rate the strength of the recommendations. Specific recommendations were rated, however, the overall guidelines were not. The Institute for Clinical Systems Improvement uses a class system: A (randomized, controlled trial), B (cohort study), C (non-randomized trial with concurrent or historical controls, case-control study, study of sensitivity and specificity of a diagnostic test, population-based descriptive study), D (cross-sectional study, case series, case report), M (meta-analysis, systematic review, decision analysis, cost-effectiveness analysis), R (consensus statement, consensus report, narrative review), X (medical opinion). The Registered Nurses’ Association of Ontario utilizes the following levels of evidence: Ia (Evidence obtained from meta-analysis or systematic review of randomized controlled trials), Ib (Evidence obtained from at least one randomized controlled trial), Ila (Evidence obtained from at least one well-designed controlled study without randomization), IIb (Evidence obtained from at least one other type of well-designed quasi-experimental study without randomization), III (Evidence obtained from well-designed non-experimental descriptive studies, such as comparative studies, correlation studies and case studies), IV (Evidence obtained from expert committee reports or opinions and/or clinical experiences of respected authorities). The Hartford Institute for Geriatric Nursing also uses a level grading system: Level I (Systematic reviews (integrative/meta-analyses) clinical practice guidelines based on systematic reviews), Level II (Single experimental study (randomized controlled trials [RCTs]), Level III (Quasi-experimental studies), Level IV (Non-experimental studies), Level V (Care report/program evaluation/narrative literature reviews), Level VI (Opinions of respected authorities/consensus panels). American Medical Directors Association guideline was developed by an interdisciplinary work group that uses a process that combined evidence- and consensus-based approaches. The type of evidence supporting the recommendations is not specifically stated. Please see attached document for further details.

1c.13 Method for rating strength of recommendation (if different from USPSTF system, also describe rating and how it relates to USPSTF): The guidelines utilize various rating systems to rate the strength of the recommendations. Specific recommendations were rated, not the overall guidelines. Please see Table 2 attached.

1c.14 Rationale for using this guideline over others: No particular guideline is recommended in this quality measure. The quality measure focuses on the outcome not on the process by which the facility reaches the outcome.

TAP/Workgroup: What are the strengths and weaknesses in relation to the subcriteria for Importance to Measure and Report? 1

Steering Committee: Was the threshold criterion, Importance to Measure and Report, met? 1

Rationale: 1

2. SCIENTIFIC ACCEPTABILITY OF MEASURE PROPERTIES

Rating: C=Completely; P=Partially; M=Minimally; N=Not at all; NA=Not applicable
Extent to which the measure, as specified, produces consistent (reliable) and credible (valid) results about the quality of care when implemented. (evaluation criteria)

2a. MEASURE SPECIFICATIONS

S.1 Do you have a web page where current detailed measure specifications can be obtained? S.2 If yes, provide web page URL:

2a. Precisely Specified

2a.1 Numerator Statement (Brief, text description of the numerator - what is being measured about the target population, e.g. target condition, event, or outcome):
The numerator is the number of short-stay residents with a discharge MDS 3.0 assessment during the selected time window who have one or more Stage 2-4 pressure ulcer(s) that are new or that have worsened on the discharge assessment compared to the previous OBRA admission or 5-day PPS assessment. Stage 1 ulcers are excluded from this measure because recent studies have identified difficulties in objectively measuring them across different populations (Lynn, 2007).

2a.2 Numerator Time Window (The time period in which cases are eligible for inclusion in the numerator):
For every quarter (3-month period), the Centers for Medicare & Medicaid Services (CMS) selects the MDS 3.0 discharge assessments from each nursing facility.

2a.3 Numerator Details (All information required to collect/calculate the numerator, including all codes, logic, and definitions):
The numerator is the number of short-stay residents with a MDS 3.0 discharge assessment during the selected time window and who have one or more Stage 2-4 pressure ulcer(s) that are new or have worsened comparing the discharge assessment (A0310.F=10, 11) and the prior OBRA admission (A0310.A=01) or the 5-day PPS assessment (A0310.B=01). On the discharge assessment, item M0800A > 0 or M0800B>0 or M0800C>0:

M0800 = Worsening in Pressure Ulcer Status Since Prior Assessment (Indicate the number of current pressure ulcers that were not present or were a lesser stage on the prior assessment: A. Stage 2, B. Stage 3, and C. Stage 4)
OR
The pressure ulcers are new or have worsened. This is indicated by comparing the discharge assessment with the prior OBRA admission or 5-day PPS assessment on item M0300 (current number of unhealed [non-epithelialized] pressure ulcers at each stage). If M0300 is equivalent or greater in the discharge assessment than in the OBRA admission or 5-day PPS assessment for each stage of ulcer, including B1 (Stage 2) OR C1 (Stage 3), or D1 (Stage 4) then they are included as having a pressure ulcer that failed to improve or is a new pressure ulcer.

Definitions of pressure ulcer stages for the MDS 3.0:
M0300 B.1 = 1 or > Stage 2: Partial thickness loss or dermis presenting as shallow open ulcer with red or pink wound bed, without slough. May also present as an intact or open/ruptured blister.
OR
M0300 C.1 = 1 or > Stage 3: Full thickness tissue loss. Subcutaneous fat may be visible but bone, tendon, or muscle is not exposed. Slough may be present but does not obscure the depth of tissue loss. May include undermining or tunneling.
OR
M0300 D.1 = 1 or > Stage 4: Full thickness tissue loss with exposed bone or tendon, or muscle. Slough or eschar may be present on some parts of the wound bed. Often includes undermining or tunneling.

2a.4 Denominator Statement (Brief, text description of the denominator - target population being measured):
All short-stay nursing facility residents except those who meet the exclusion criteria.

2a.5 Target population gender: Female, Male

Comment [KP8]: 2a. The measure is well defined and precisely specified so that it can be implemented consistently within and across organizations and allow for comparability. The required data elements are of high quality as defined by NQF’s Health Information Technology Expert Panel (HITEP).
<table>
<thead>
<tr>
<th>Section</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>2a.6</td>
<td><strong>Target population age range:</strong> The target population includes people of all ages who are short-stay residents in the nursing facility.</td>
</tr>
<tr>
<td>2a.7</td>
<td><strong>Denominator Time Window</strong> <em>(The time period in which cases are eligible for inclusion in the denominator):</em> For every quarter (3-month period), CMS selects the MDS 3.0 discharge assessments from each nursing facility.</td>
</tr>
<tr>
<td>2a.8</td>
<td><strong>Denominator Details</strong> <em>(All information required to collect/calculate the denominator - the target population being measured - including all codes, logic, and definitions):</em> The denominator is the number of short-stay residents who have been assessed with MDS 3.0 discharge assessments during the selected time window and whose date of discharge is less than or equal to 100 days since their most recent entry date (A1600) for the OBRA admission or 5-day PPS assessment, except for those meeting the exclusion criteria.</td>
</tr>
<tr>
<td>2a.9</td>
<td><strong>Denominator Exclusions</strong> <em>(Brief text description of exclusions from the target population):</em> A short-stay resident is excluded from the denominator if there is no discharge assessment or if missing data precludes calculation of the measure. Short-stay facilities are excluded from public reporting if they have fewer than 20 residents due to small sample size.</td>
</tr>
<tr>
<td>2a.10</td>
<td><strong>Denominator Exclusion Details</strong> <em>(All information required to collect exclusions to the denominator, including all codes, logic, and definitions):</em> A0310F = 10 discharge assessment (return not anticipated) or 11 discharge assessment (return anticipated) = missing OR M0800 = Worsening in Pressure Ulcer Status Since Prior Assessment (Indicate the number of current pressure ulcers that were not present or were a lesser stage on the prior assessment: A. Stage 2, B. Stage 3, and C. Stage 4) = missing OR if any of the following: M0800A. (Stage 2), B. (Stage 3), or C. (Stage 4) are not completed and any of the values in the completed item(s) = 0, then the missing data preclude calculation of the measure and so are excluded. Short-stay facilities with fewer than 20 residents are excluded from public reporting because of small sample size.</td>
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<tr>
<td>2a.11</td>
<td><strong>Stratification Details/Variables</strong> <em>(All information required to stratify the measure including the stratification variables, all codes, logic, and definitions):</em> This is not applicable.</td>
</tr>
<tr>
<td>2a.12-13</td>
<td><strong>Risk Adjustment Type:</strong> Case-mix adjustment</td>
</tr>
<tr>
<td>2a.14</td>
<td><strong>Risk Adjustment Methodology/Variables</strong> <em>(List risk adjustment variables and describe conceptual models, statistical models, or other aspects of model or method):</em> Resident-level limited covariate risk adjustment is performed. Covariates are based on the 5-day PPS assessment and include residents who have healed pressure ulcer(s), require limited or more assistance in bed, have bowel incontinence at least once a week, diabetes or peripheral vascular disease, or low Body Mass Index (BMI between 12 -19). Resident-level covariates are used in a logistic regression model to calculate a resident-level expected QM score (the probability that the resident will evidence the outcome, given the presence or absence of characteristics measured by the covariates). Then, an average of all resident-level expected QM score for the nursing facility is calculated to create a facility-level expected QM score. The final facility-level adjusted QM score is based on a calculation which combines the facility-level expected score and the facility-level observed score.</td>
</tr>
<tr>
<td>2a.15-17</td>
<td><strong>Detailed risk model available Web page URL or attachment:</strong> URL</td>
</tr>
</tbody>
</table>

Rating: C=Completely; P=Partially; M=Minimally; N=Not at all; NA=Not applicable
Type of Score: Ratio

Calculation Algorithm (Describe the calculation of the measure as a flowchart or series of steps):
For each facility, the number of short-stay residents meeting the numerator criteria and the number of (non-excluded) residents meeting the denominator criteria are counted. These numbers are also counted for the covariate measure, which are residents who have healed pressure ulcer(s), require limited or more assistance in bed, have bowel incontinence at least one/week, diabetes or peripheral vascular disease, or low Body Mass Index (BMI) (between 12-19) as reported on the resident’s 5-day PPS assessment.

The covariate scores are entered into a logistic regression equation and the result is an expected score for the resident for that quality measure. The logistic regression equations are of the form: where e is the base of natural logarithms and x is a linear combination of the logistic regression coefficients and the covariate scores of the form:

\[ C_0 + C_1 \cdot COVA + C_2 \cdot COVB + \ldots \]

where \( C_0 \) is the logistic regression constant, \( C_1 \) is the logistic regression coefficient for the first covariate (where applicable), \( COVA \) is the resident-level score for the first covariate, \( C_2 \) is the logistic regression coefficient for the second covariate, and \( COVB \) is the resident-level score for the second covariate (where applicable), etc. The regression constant and regression coefficients are numbers obtained through statistical logistic regression analysis.

The expected score for the measure is then calculated as the expected number of residents in the facility meeting the numerator criteria divided by all non-excluded residents in the denominator. (1)


Describe the method for discriminating performance (e.g., significance testing):
Because the computed scores are not estimates, but include all residents who meet the measure criteria, in terms of discriminating performance, the computed scores can be used to make valid comparisons.

Sampling (Survey) Methodology If measure is based on a sample (or survey), provide instructions for obtaining the sample, conducting the survey and guidance on minimum sample size (response rate):
This is not applicable.

Data Source (Check the source(s) for which the measure is specified and tested)
Electronic clinical data

Data source/data collection instrument (Identify the specific data source/data collection instrument, e.g. name of database, clinical registry, collection instrument, etc.):
Nursing Home MDS 3.0

Data source/data collection instrument reference web page URL or attachment: URL
http://www.cms.hhs.gov/NursingHomeQualityInit/25_NHQIMDS30.asp#TopOfPage

Data dictionary/code table web page URL or attachment: URL
http://www.cms.hhs.gov/NursingHomeQualityInit/25_NHQIMDS30.asp#TopOfPage

Level of Measurement/Analysis (Check the level(s) for which the measure is specified and tested)
Population: national, Facility/Agency

Care Settings (Check the setting(s) for which the measure is specified and tested)
Nursing home (NH) / Skilled Nursing Facility (SNF)

Clinical Services (Healthcare services being measured, check all that apply)

Comment [KP10]: Reliability testing demonstrates the measure results are repeatable, producing the same results a high proportion of the time when assessed in the same population in the same time period.
Three major tests of the reliability of the short-stay pressure ulcer quality measure have been conducted. First, the MDS 2.0 measure items and selected existing quality measure were tested in the Data Assessment and Verification (DAVE 2) project conducted by Abt Associates. This project used a nationwide sample of randomly selected nursing homes using MDS assessments for the period April 1 to December 31, 2006. DAVE 2 performed 173 two-stage reviews.

Second, the University of Colorado used two sources of data: national facility-level quality measure data from 2003 Quarter 3 (Q3) through 2006 Q3 came from the Quality Improvement and Evaluation System (QIES) MDS Express Reports on the CMS intranet; Online Survey, Certification, and Reporting (OSCAR) data related to facility characteristics (e.g., state, resident census, number of beds, staffing) and certification survey results were downloaded from QIES Workbench. A 10% random sample of all Medicare-certified nursing facilities was also downloaded from MDS assessment records. Analyses were based on complete MDS data from January 2005 through March 2006, nearly complete data for April 2006, and partial data for May and June 2006.

Third, testing of the reliability of MDS 3.0 data items underlying the pressure ulcer quality measure and a comparison with the MDS 2.0 quality measures were conducted by RAND as part of the MDS 3.0 development process. A representative sample of for-profit and not-for-profit facilities and hospital-based and freestanding facilities was recruited for the study, which included 71 community nursing homes in 8 states, 19 VA nursing homes, and 1,402 nursing home residents for the pressure ulcer quality measure.


### Analytic Method

**Description of Data/Sample and Size:**
Three sets of analytic methods were used. First, in the DAVE 2 Project, trained nurse reviewers selected a current resident with a recent assessment performed by the nursing home (NH) within the last 14 days. In the first stage of this review, the nurse reviewer conducted a blind reassessment of the resident using standard MDS assessment and coding procedures (e.g., examination of the medical record; observation of the resident; interview of staff, resident, and family; and use of coding criteria). In the second stage of this assessment, the DAVE 2 nurse reviewer’s assessment was compared with the corresponding nursing home assessment, and each discrepancy was reconciled, with the nursing home assessor and the nurse reviewer agreeing on the appropriate response. In addition to data entering the facility MDS code, the DAVE 2 code, and the reconciled code into the MDS-QC data entry software, the DAVE 2 nurse reviewer entered a “reason code” to attribute the cause of the discrepancy, per MDS item reviewed, to an established list of reasons.

Second, in terms of measure stability, which is not exactly the same as reliability but is a concept related to it, the University of Colorado examined the percentage of facilities that had a change in ranking from one quarter to the next of at least 3 deciles. This indicator of stability was computed for each of the 12 pairs of adjacent quarters for which data were available (2003 Q3 through 2006 Q3). The range of stability measures across the 12 comparisons was very small (i.e., the difference between the maximum and minimum values), indicating that measure stability is quite constant over time. For pressure ulcers, the minimum percentage was 27.0%, and the maximum percentage was 29.0%.

Third, the national test of MDS 3.0 items examined agreement between assessors (reliability). QIOs were employed to identify gold-standard (research) nurses and recruit community nursing facilities to participate in the national evaluation. The gold-standard nurses were trained in the MDS 3.0 instrument and, in turn, trained a facility nurse from each participating nursing facility in their home states. Quality measures using the MDS 2.0 and the MDS 3.0 were calculated and then compared, with correlations and kappas calculated.

1. Abt Associates, Inc.; Stepwise Systems, Inc.; Qualidigm. Data Assessment and Verification (DAVE 2)
2b.3 Testing Results (reliability statistics, assessment of adequacy in the context of norms for the test conducted):  
As part of the DAVE 2 project, Abt Associates used two methods to assess the reliability of the MDS 2.0 quality measures. (1) First, for each MDS data element, the rate of discrepancies between the reconciled and original facility assessments has been reported. For the pressure ulcer item, M26a the two-stage review discrepancy rate was 6.0%. Second, Abt reported the rate of discrepancies between selected quality measures, computed from facility data, and its counterpart, computed from reconciled data. Data for the current quality measure was not analyzed during that study. However, the University of Colorado researchers noted that the low discrepancy rate across providers should lead to a low measure level discrepancy rate as well. (2)

Second, in terms of measure stability, the University of Colorado examined the percentage of facilities that had a change in ranking of at least three deciles from one quarter to the next. (2) For short-stay pressure ulcer, 21.5% of facilities had a change of 3 deciles or more from one quarter to the next.

Third, in their testing of the MDS 3.0, RAND compared the results on the nursing home quality measures using the MDS 3.0 and the MDS 2.0, both at the individual resident level and at the facility level. (3) At the resident level, the pressure ulcer rate using the MDS 2.0 was 25.1%; using the MDS 3.0 it was 23.6%; the Kappa was 0.92, the correlation was 0.92, and the percent agreement was 97.0. Kappa is a statistical measure of inter-rater agreement for qualitative data, ranging from 0.0 to 1.0. A rating of 0.70 is considered “substantial agreement.” At the facility level, the MDS 2.0 rate of pressure ulcers was 26.9% and the MDS 3.0 rate was 26.0%, with a correlation of 0.98, which is quite high. However, the researchers analyzed the prevalence versus incidence (actual quality measure).

2c. Validity testing

2c.1 Data/sample (description of data/sample and size): The data came from two sources: national facility-level pressure measure data from 2003 Q3 through 2006 Q3 came from the QIES MDS Express Reports on the CMS intranet; OSCAR data related to facility characteristics (e.g., state, resident census, number of beds, staffing) and certification survey results were downloaded from QIES Workbench. A 10% random sample of all Medicare-certified nursing facilities was also downloaded from MDS assessment records. Analyses were based on complete MDS data from January 2005 through March 2006, as well as nearly complete data for April 2006 and partial data for May and June 2006.


2c.2 Analytic Method (type of validity & rationale, method for testing): Yes, the testing is incomplete because the reliability testing for the measure is based on the MDS 2.0. The underlying pressure ulcer items have significantly changed from the MDS 2.0 to the MDS 3.0 although RAND did...
The analysis evaluated measure validity in a number of ways: 1) examining the expected positive influence of public reporting on quality of care, 2) assessing the degree to which pressure ulcer rates have improved over time; 3) evaluating convergent validity, an assessment of the correlation of the quality measure with all other measures; 4) determining if the quality measure rate was influenced by factors that are unrelated to facility quality, and 5) evaluating seasonal variations in pressure ulcer rates across the 13 quarters of data. The analysis also computed descriptive statistics and conducted a one-way analysis of variance (ANOVA) for the measure to examine the amount of variance in pressure ulcer rates explained in the state in which a facility was located.

2c.3 Testing Results (statistical results, assessment of adequacy in the context of norms for the test conducted):

To evaluate convergent validity, an assessment of the correlation of the current MDS 2.0-based quality measure with all other measures was performed. A correlation of approximately 0.32 indicates that one measure explains about 10% of the variance in the other measure. Correlations smaller than .30, therefore, indicate a weak relationship between two measures. One would expect a clinically related quality measure, such as pressure ulcers, to be correlated. Overall, the pressure ulcer measure was not well correlated to other quality measures except high risk pressure ulcer for long-stay residents.

See attached Table 2: Correlations of Quality Measures—Excluding Vaccination Quality Measures (Facility Level).

The analysis found that public reporting appears to have had some influence on the decreased pressure ulcers over time, for the current MDS 2.0-based quality measure as evidenced by the decline in the triggering rate from 20.4 to 17.1. However, the quality measure did demonstrate seasonal variation, suggesting that the measure is influenced by factors beyond the provision of care.

See attached Table 3: Measure Trends Over Time.

Also, there was variability across facilities in the rates of pressure ulcers for short-stay residents, suggesting that it is possible for facilities to improve. As presented in the table below, the national overall nursing facility mean in 2006 Q3 was 17.3, and the standard deviation was 9.3. The short-stay pressure ulcer quality measure demonstrated variability across facilities; from 7.0% at the 10th percentile to 28.7% at the 90th percentile, with only 1.1% of facilities reporting no residents with pressure ulcers.

See attached Table 1: Measure Variability Across Facilities.

The most recent state and national averages for the current MDS 2.0 pressure ulcer quality measure are reported on CMS’s Nursing Home Compare Web site for the target quarter ending in June 2009. The data continue to demonstrate the ongoing gap in facility performance; the national average was 16.9%, and state averages ranged from a low of 11.3% to a high of 25.6%.

The statistical limitations of this measure may reflect a limited clinical relationship of pressure ulcers to the other quality measures, and while the variation in rate among states makes it difficult to compare facilities between different states, the measure remains a valuable guide between facilities within the same state.

2d. Exclusions Justified

2d.1 Summary of Evidence supporting exclusion(s):
All assessments of short-stay residents for whom complete data are available are included.

2d.2 Citations for Evidence:
This is not applicable.

Comment [KP14]: 2d. Clinically necessary measure exclusions are identified and must be:
• supported by evidence of sufficient frequency of occurrence so that results are distorted without the exclusion;
AND
• a clinically appropriate exception (e.g., contraindication) to eligibility for the measure focus;
AND
• precisely defined and specified:
  • if there is substantial variability in exclusions across providers, the measure is specified so that exclusions are computable and the effect on the measure is transparent (i.e., impact clearly delineated, such as number of cases excluded, exclusion rates by type of exclusion);
  • if patient preference (e.g., informed decision-making) is a basis for exclusion, there must be evidence that it strongly impacts performance on the measure and the measure must be specified so that the information about patient preference and the effect on the measure is transparent (e.g., numerator category computed separately, denominator exclusion category computed separately).

Comment [KP15]: 10 Examples of evidence that an exclusion distorts measure results include, but are not limited to: frequency of occurrence, sensitivity analyses with and without the exclusion, and variability of exclusions across providers.
2d.3 Data/sample (description of data/sample and size): This is not applicable.

2d.4 Analytic Method (type analysis & rationale): This is not applicable.

2d.5 Testing Results (e.g., frequency, variability, sensitivity analyses): This is not applicable.

2e. Risk Adjustment for Outcomes/Resource Use Measures

2e.1 Data/sample (description of data/sample and size): Samples for two “target periods” were drawn:
1. “Current Period” target sample for computing quality measures: all U.S. nursing facilities and residents selected from the target quarter.
2. “Prior Year” target sample for estimating logistic regressions: post-acute care (PAC) patients from a 20 percent random sample of all nursing facilities with a post-acute care admission in the period 2001 Q4 through 2002 Q3.
All PAC patient records included, for each target period:
1. A discharge skilled nursing facility (SNF) assessment (most recent) in the target period.
2. A 5-day SNF PPS assessment from the same stay, if available.
3. A recent admission assessment, if available:
   a. A target assessment (most recent)
   b. A prior assessment preceding the target assessment, if available
   c. A most recent full assessment, if available
The information for risk adjustment is from:

2e.2 Analytic Method (type of risk adjustment, analysis, & rationale):
The approach involves using logistic regression to adjust quality measure scores directly. This method of adjustment uses resident-level covariates that have been found to increase the risks of an outcome.
First, resident-level covariates were used in a logistic regression model to calculate a resident-level expected quality measure score (the probability that the resident will evidence the outcome, given the presence or absence of characteristics measured by the covariates).
Then, an average of all resident-level expected quality measure scores for the nursing facility was calculated to create a facility-level expected quality measure score.
The final facility-level adjusted quality measure score was based on a calculation that combines the facility-level expected score and the facility-level observed score.

2e.3 Testing Results (risk model performance metrics):
A review by the University of Colorado used data from two sources: national facility-level quality measure data from 2003 Q3 through of 2006 Q3, which came from the QIES MDS Express Reports on the CMS Intranet; and OSCAR data related to facility characteristics (e.g., state, resident census, number of beds, staffing) and certification survey results were downloaded from the QIES Workbench. A 10% random sample of all Medicare-certified nursing facilities was also downloaded from MDS assessment records. Analyses were based on complete MDS data from January 2005 through March 2006, nearly complete data for April 2006, and partial data for May and June 2006. The analysis evaluated the risk adjustment model at the resident level and the facility level, generating R-square and C statistics. R-square indicates the proportion of the variance in measure performance that is accounted for by the covariates. The C-statistic gives the percentage of the time the observed value and the expected value move in the same direction, ranging from 0.5 (poor, no better than chance) to 1.0 (observed and predicted always move in the same direction). The analysis found an R2 of .65 and a C of .67; neither statistic met the threshold for predictive adequacy (0.10 for R-square and 0.70 for C).

2e.4 If outcome or resource use measure is not risk adjusted, provide rationale: This is not applicable.

Comment [KP16]: 2e. For outcome measures and other measures (e.g., resource use) when indicated:
• an evidence-based risk-adjustment strategy (e.g., risk models, risk stratification) is specified and is based on patient clinical factors that influence the measured outcome (but not disparities in care) and are present at start of care. [Strict Benchmark not defined. OR rationale/data support no risk adjustment.]

Comment [k17]: 13 Risk models should not obscure disparities in care for populations by including factors that are associated with differences/inequalities in care such as race, socioeconomic status, gender (e.g., poorer treatment outcomes of African American men with prostate cancer, inequalities in treatment for CVD risk factors between men and women). It is preferable to stratify measures by race and socioeconomic status rather than adjusting out differences.
2f. Identification of Meaningful Differences in Performance

2f.1 Data/sample from Testing or Current Use (description of data/sample and size): Yes, the testing is based on MDS 2.0 items.

An analytical team at the University of Colorado Health Sciences Center examined the short-stay pressure ulcer rates at the facility level based on the current measure and underlying 2.0 items.(1) For 10,056 facilities, the mean for the pressure ulcer measure was 17.1% and the standard deviation was 9.3%. The quality measure varied from 7.0% at the 10th percentile to 28.7% at the 90th percentile; only 1.1% of facilities had no residents with pressure ulcers.

See attached Table 1: Measure Variability Across Facilities.

The most recent state and national averages for the current pressure ulcer quality measure are reported on CMS's Nursing Home Compare Web site for the target quarter ending in June 2009.(2) The data continue to demonstrate the ongoing gap in facility performance; the national average was 10% and state ranged from a low of 8.4% to a high of 18.7%.(2)


2f.2 Methods to identify statistically significant and practically/meaningfully differences in performance (type of analysis & rationale): Because the computed scores are not estimates, but include all residents who meet the measure criteria, in terms of discriminating performance, the computed scores can be used to make valid comparisons.

2f.3 Provide Measure Scores from Testing or Current Use (description of scores, e.g., distribution by quartile, mean, median, SD, etc.; identification of statistically significant and meaningfully differences in performance):

For 10,056 facilities, the mean pressure ulcer rate was 17.1 percent with a standard deviation of 9.3 percent. The attached Table 1: Measure Variability Across Facilities reports the full result of the analysis.

2g. Comparability of Multiple Data Sources/Methods

2g.1 Data/sample (description of data/sample and size): This is not applicable.

2g.2 Analytic Method (type of analysis & rationale):

This is not applicable.

2g.3 Testing Results (e.g., correlation statistics, comparison of rankings):

This is not applicable.

2h. Disparities in Care

2h.1 If measure is stratified, provide stratified results (scores by stratified categories/cohorts): This is not applicable.

2h.2 If disparities have been reported/identified, but measure is not specified to detect disparities, provide follow-up plans:

This is not applicable; however, RTI will analyze the proposed quality measure data when MDS 3.0 is implemented.
### 3. USABILITY

**Extent to which intended audiences (e.g., consumers, purchasers, providers, policy makers) can understand the results of the measure and are likely to find them useful for decision making.** (evaluation criteria)

### 3a. Meaningful, Understandable, and Useful Information

#### 3a.1 Current Use: In use

#### 3a.2 Use in a public reporting initiative (disclosure of performance results to the public at large) (If used in a public reporting initiative, provide name of initiative(s), locations, Web page URL(s). If not publicly reported, state the plans to achieve public reporting within 3 years):


#### 3a.3 If used in other programs/initiatives (If used in quality improvement or other programs/initiatives, name of initiative(s), locations, Web page URL(s). If not used for QI, state the plans to achieve use for QI within 3 years):

*CMS expects that the short-stay quality measure will be used by nursing facilities as a tool to improve quality of care by preventing pressure ulcers; when pressure ulcers do occur, nursing facilities can use the measure to ensure that they are treated in a timely fashion and according to guidelines. Data on facility performance on the quality measures are also used by surveyors to identify problem areas when they inspect nursing homes.*

#### Testing of Interpretability (Testing that demonstrates the results are understood by the potential users for public reporting and quality improvement)

#### 3a.4 Data/sample (description of data/sample and size): A recent study by Castle found that consumers could accurately interpret the quality information given for all the current nursing facility quality measures based on MDS 2.0 reported by Nursing Home Compare.(1)

In the Castle article, an initial sample of 8,000 family members with elders living in one of 200 randomly selected nursing facilities was used.(1) In each facility, one family member (or significant other) was identified as the family contact person for each of 40 residents by nursing facility staff. A total of 615 facilities were approached before the target of 200 participating facilities was achieved, giving a facility participation rate of 33%. From these 200 facilities, a total of 4,754 surveys were returned (i.e., family response rate = 59%).


#### 3a.5 Methods (e.g., focus group, survey, QI project):

A comprehension index was developed to examine whether the information contained in Nursing Home Compare for each currently reported quality measure (based on MDS 2.0) was understood by family members.(1) The measures ranged from 0.0 to 8.0.


#### 3a.6 Results (qualitative and/or quantitative results and conclusions):

Castle found that 31% of the consumers used the Internet in choosing a nursing facility; 12% recalled using Nursing Home Compare, and in general, the consumers' comprehension index scores indicated a relatively good understanding of the measures.(1) The comprehension index for the current pressure ulcer measure,

3b/3c. Relation to other NQF-endorsed measures

3b.1 NQF # and Title of similar or related measures:

(for NQF staff use) Notes on similar/related endorsed or submitted measures:

3b. Harmonization
If this measure is related to measure(s) already endorsed by NQF (e.g., same topic, but different target population-setting/data source or different topic but same target population):

3b.2 Are the measure specifications harmonized? If not, why?
The measure specifications are not harmonized with any other pressure ulcer measures but the underlying MDS 3.0 items that make up the proposed measure move toward that goal. The NQF’s initiative “National Voluntary Consensus Standards for Developing a Framework for Measuring Quality for Prevention and Management of Pressure Ulcers,” begun in 2008, has several objectives, including determining how to measure the incidence, prevalence, and staging of pressure ulcers and harmonizing measure specifications across settings of care. The revised approach to documenting pressure ulcers in the MDS 3.0 is part of that process.

The current publicly reported nursing facility quality measure reports on Stage 1-4 pressure ulcers that are new or that have not improved based on the MDS 2.0 for short-stay residents. The MDS 2.0 items for pressure ulcers were problematic for wound care experts because they (1) used reverse staging, which does not reflect the pathophysiology of wound healing; (2) failed to capture size or change in size and therefore missed improvement; (3) inappropriately “staged” stasis ulcers; (4) failed to document pressure ulcers that were present on admission, and (5) did not allow for the category “unstageable.” The goal of the MDS 3.0 pressure ulcer items was to align MDS 3.0 with accepted best practices.

The proposed measure reports Stage 2-4 pressure ulcers based on new items on the MDS 3.0. As Saliba and Buchanan noted during the development of the MDS 3.0, whenever possible, they included items or language used in other health care settings in order to improve communication across settings and providers (e.g., the pressure ulcer items included in the National Pressure Ulcer Advisory Panel’s PUSH tool are used to describe pressure ulcers in the MDS 3.0). (1)

The variation across populations and measure specification of the range of NQF endorsed pressure ulcer measures will make harmonization challenging. The other NQF-endorsed pressure ulcer measures vary by examining only hospital-acquired (nosocomial) Stage II or greater pressure ulcers on the day of the prevalence study or by focusing on the home care setting and whether (1) patients with assessed risk for pressure ulcers have a physician-ordered plan of care that includes intervention(s) to prevent them, (2) patients with assessed risk for pressure ulcers have interventions for pressure ulcer prevention that were implemented during their episode of home care, or (3) patients were assessed for risk of pressure ulcers at start/resumption of home health care.

3c. Distinctive or Additive Value

3c.1 Describe the distinctive, improved, or additive value this measure provides to existing NQF-endorsed measures:

The proposed measure reports Stage 2-4 pressure ulcers based on new items on the MDS 3.0. As Saliba and Buchanan noted during the development of the MDS 3.0, whenever possible, they included items or language used in other health care settings in order to improve communication across settings and providers (e.g., the pressure ulcer items included in the National Pressure Ulcer Advisory Panel’s PUSH tool are used to describe pressure ulcers in the MDS 3.0). (1) Therefore, the proposed measure based on the new MDS 3.0 pressure ulcer items better aligns the measure with accepted best practices. Additionally, Stage 1 ulcers are not included in this measure because recent studies identified difficulties in objectively measuring them across different populations (Lynn, 2007)

The other NQF-endorsed pressure ulcer measures differ from this because they only examine hospital acquired (nosocomial), Stage 2 or worse pressure ulcers, on the day of the prevalence study. Or they focus on the home care setting and whether: (1) patients with assessed risk for pressure ulcers have a physician-ordered plan of care that includes prevention intervention(s), (2) patients with assessed risk for pressure ulcers have interventions for pressure ulcer prevention that were implemented during their episode of home care, or (3) patients were assessed for risk of pressure ulcers at start/resumption of home health care.


5.1 If this measure is similar to measure(s) already endorsed by NQF (i.e., on the same topic and the same target population), Describe why it is a more valid or efficient way to measure quality:

TAP/Workgroup: What are the strengths and weaknesses in relation to the subcriteria for Usability?

Steering Committee: Overall, to what extent was the criterion, Usability, met?

Rationale:

4. FEASIBILITY

Extent to which the required data are readily available, retrievable without undue burden, and can be implemented for performance measurement. (evaluation criteria)

4a. Data Generated as a Byproduct of Care Processes

4a.1-2 How are the data elements that are needed to compute measure scores generated?

Data generated as byproduct of care processes during care delivery (Data are generated and used by healthcare personnel during the provision of care, e.g., blood pressure, lab value, medical condition), Coding/abstraction performed by someone other than person obtaining original information (E.g., DRG, ICD-9 codes on claims, chart abstraction for quality measure or registry)

4b. Electronic Sources

4b.1 Are all the data elements available electronically? (elements that are needed to compute measure scores are in defined, computer-readable fields, e.g., electronic health record, electronic claims)

No

4b.2 If not, specify the near-term path to achieve electronic capture by most providers.

Not applicable.

4c. Exclusions

Comment [KP25]: 3c. Review of existing endorsed measures and measure sets demonstrates that the measure provides a distinctive or additive value to existing NQF-endorsed measures (e.g., provides a more complete picture of quality for a particular condition or aspect of healthcare, is a more valid or efficient way to measure).

Comment [KP26]: 4a. For clinical measures, required data elements are routinely generated concurrent with and as a byproduct of care processes during care delivery. (e.g., BP recorded in the electronic record, not abstracted from the record later by other personnel; patient self-assessment tools, e.g., depression scale; lab values, meds, etc.)

Comment [KP27]: 4b. The required data elements are available in electronic sources. If the required data are not in existing electronic sources, a credible, near-term path to electronic collection by most providers is specified and clinical data elements are specified for transition to the electronic health record.

Comment [KP28]: 4c. Exclusions should not require additional data sources beyond what is required for scoring the measure (e.g., numerator and denominator) unless justified as supporting measure validity.
4c.1 Do the specified exclusions require additional data sources beyond what is required for the numerator and denominator specifications?
No

4c.2 If yes, provide justification.

4d. Susceptibility to Inaccuracies, Errors, or Unintended Consequences

4d.1 Identify susceptibility to inaccuracies, errors, or unintended consequences of the measure and describe how these potential problems could be audited. If audited, provide results.
The proposed measure differs from the current measure because of improvements in the underlying items and the elimination of Stage 1 pressure ulcers from reporting. The goal of the new MDS 3.0 items was to align the pressure ulcer items with accepted best practices using the National Pressure Ulcer Advisory Panel’s PUSH tool to describe pressure ulcers. In their validation work, Saliba and Buchanan reported that according to staff feedback, “89% felt that definitions were clear (3% disagreed).”(1)

For the updated MDS 3.0 pressure ulcer items, average gold-standard to gold-standard kappa was .905. Average gold-standard to facility-nurse kappa was .937. However, implementation on a nationwide basis does not begin until October 2010, and the data will be analyzed to address any concerns regarding inaccuracies, errors, or unintended consequences.


4e. Data Collection Strategy/Implementation

4e.1 Describe what you have learned/modified as a result of testing and/or operational use of the measure regarding data collection, availability of data/missing data, timing/frequency of data collection, patient confidentiality, time/cost of data collection, other feasibility/implementation issues:
The data collection method, the MDS, is already in operation and has been for many years.

4e.2 Costs to implement the measure (costs of data collection, fees associated with proprietary measures):
The data are collected as part of an existing, legally mandated process. There will be no additional costs to collect this information because it is already collected. However, the pressure ulcer measure items have changed substantially in the MDS 3.0 and require additional training for facility staff.

4e.3 Evidence for costs:
This is not applicable.

4e.4 Business case documentation: The proposed measure relies on data from the MDS 3.0. As there is no change in the data collection method for the MDS 3.0 as compared with its predecessor, the MDS 2.0, we do not anticipate any additional burden to nursing facilities. MDS 2.0, and soon to be MDS 3.0, data are collected as part of an existing, federally mandated process used for payment and quality monitoring purposes.

TAP/Workgroup: What are the strengths and weaknesses in relation to the subcriteria for Feasibility?

Steering Committee: Overall, to what extent was the criterion, Feasibility, met?
Rationale:

RECOMMENDATION

(for NQF staff use) Check if measure is untested and only eligible for time-limited endorsement.
<table>
<thead>
<tr>
<th>CONTACT INFORMATION</th>
</tr>
</thead>
<tbody>
<tr>
<td>Co.1 Measure Steward (Intellectual Property Owner)</td>
</tr>
<tr>
<td>Co.1 Organization</td>
</tr>
<tr>
<td>Centers for Medicare &amp; Medicaid Services, 7500 Security Boulevard, Mail Stop S3-02-01, Baltimore, Maryland, 21244-1850</td>
</tr>
<tr>
<td>Co.2 Point of Contact</td>
</tr>
<tr>
<td>Judith, Tobin, PT, MBA, <a href="mailto:Judith.Tobin@cms.hhs.gov">Judith.Tobin@cms.hhs.gov</a>, 410-786-6892-</td>
</tr>
<tr>
<td>Measure Developer If different from Measure Steward</td>
</tr>
<tr>
<td>Co.3 Organization</td>
</tr>
<tr>
<td>RTI International, 1440 Main Street, Suite 310, Waltham, Massachusetts, 02451-1623</td>
</tr>
<tr>
<td>Co.4 Point of Contact</td>
</tr>
<tr>
<td>Roberta, Constantine, RN, MBA, PhD, <a href="mailto:rconstantine@rti.org">rconstantine@rti.org</a>, 781-434-1711-</td>
</tr>
<tr>
<td>Co.5 Submitter If different from Measure Steward POC</td>
</tr>
<tr>
<td>Roberta, Constantine, RN, MBA, PhD, <a href="mailto:rconstantine@rti.org">rconstantine@rti.org</a>, 781-434-1711-, RTI International</td>
</tr>
<tr>
<td>Co.6 Additional organizations that sponsored/participated in measure development</td>
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<table>
<thead>
<tr>
<th>ADDITIONAL INFORMATION</th>
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<tbody>
<tr>
<td>Workgroup/Expert Panel involved in measure development</td>
</tr>
<tr>
<td>Ad.1 Provide a list of sponsoring organizations and workgroup/panel members’ names and organizations. Describe the members’ role in measure development.</td>
</tr>
<tr>
<td>See attached Table 4. Nursing Home Quality Measures Technical Expert Panel (January 2009) showing a list of workgroup or panel member names and organizations.</td>
</tr>
<tr>
<td>This technical expert panel met during 2 days in January 2009 to review the environmental scan of the current quality measures and make recommendations regarding their transition from MDS 2.0 to MDS 3.0.</td>
</tr>
<tr>
<td>Ad.2 If adapted, provide name of original measure: This measure was adapted from the measure of the same name derived from MDS 2.0 data.</td>
</tr>
<tr>
<td>Ad.3-5 If adapted, provide original specifications URL or attachment</td>
</tr>
<tr>
<td>Measure Developer/Steward Updates and Ongoing Maintenance</td>
</tr>
<tr>
<td>Ad.6 Year the measure was first released: 2002</td>
</tr>
<tr>
<td>Ad.7 Month and Year of most recent revision: 02, 2010</td>
</tr>
<tr>
<td>Ad.8 What is your frequency for review/update of this measure? Every 3 years</td>
</tr>
<tr>
<td>Ad.9 When is the next scheduled review/update for this measure? 02, 2013</td>
</tr>
<tr>
<td>Ad.10 Copyright statement/disclaimers:</td>
</tr>
<tr>
<td>Ad.11 -13 Additional Information web page URL or attachment: Attachment Pressure Ulcers Short Stay tables_FINAL-634045018897767500.doc</td>
</tr>
<tr>
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