# Proposed Measure Specifications for Measures Proposed in the FY 2017 SNF QRP NPRM

# Prepared for

**Center for Clinical Standards and Quality** 

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# PROPOSED MEASURE SPECIFICATIONS FOR MEASURES PROPOSED IN THE FY 2017 SNF QRP NPRM

# **RTI** International

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# SECTION 1 CROSS-SETTING MEASURES DEVELOPMENT WORK: AN INTRODUCTION

The Improving Medicare Post-Acute Care Transformation Act of 2014 (IMPACT Act), enacted Oct. 6, 2014, directs the Secretary of Health and Human Services to "specify quality measures on which Post-Acute Care (PAC) providers are required under the applicable reporting provisions to submit standardized patient assessment data" in several domains, including medication reconciliation and resource use measures, including Medicare spending per beneficiary, discharge to community and all-condition risk-adjusted potentially preventable readmission rates. The IMPACT Act requires the implementation of measures to address these measure domains in home health agencies (HHAs), skilled nursing facilities (SNFs), long-term care hospitals (LTCHs), and inpatient rehabilitation facilities (IRFs).

The IMPACT Act also requires, to the extent possible, the submission of such quality measure data through the use of a PAC assessment instrument and the modification of such instrument as necessary to enable such use; for SNFs, the Minimum Data Set (MDS) 3.0 will be used.

For more information on the statutory history of the SNF QRP, please refer to the FY 2016 SNF PPS final rule at <a href="https://www.gpo.gov/fdsys/pkg/FR-2015-08-04/pdf/2015-18950.pdf">https://www.gpo.gov/fdsys/pkg/FR-2015-08-04/pdf/2015-18950.pdf</a>. More information on the IMPACT Act is available at <a href="https://www.govtrack.us/congress/bills/113/hr4994">https://www.govtrack.us/congress/bills/113/hr4994</a>.

In this document, we present specifications for the following three (3) measures proposed for adoption for the SNF QRP through the FY 2017 SNF PPS Final Rule:

- 1. Discharge to Community- Post Acute Care (PAC) Skilled Nursing Facility Quality Reporting Program;
- 2. Potentially Preventable 30-Days Post-Discharge Readmission Measure for Skilled Nursing Facility Quality Reporting Program;
- 3. Drug Regimen Review Conducted with Follow-Up for Identified Issues- Post Acute Care (PAC) Skilled Nursing Facility Quality Reporting Program.

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# SECTION 2 MEASURES AFFECTING THE FY 2018 PAYMENT DETERMINATION AND SUBSEQUENT YEARS

# 2.1 Discharge to Community-Post Acute Care (PAC) Skilled Nursing Facility (SNF) Quality Reporting Program (QRP)

# 2.1.1 Measure Description

Sections 1899B(d)(1)(B) and 1899B(a)(2)(E)(ii) of the Act require the Secretary to specify a measure to address the resource use and other measures domain of discharge to community by SNFs, LTCHs, and IRFs by October 1, 2016, and HHAs by January 1, 2017. We are proposing to adopt the measure, Discharge to Community-Post Acute Care (PAC) Skilled Nursing Facility (SNF) Quality Reporting Program (QRP) for the SNF QRP as a Medicare FFS claims-based measure to meet this requirement.

This proposed measure assesses successful discharge to the community from a PAC setting, with successful discharge to the community including no unplanned rehospitalizations and no death in the 31 days following discharge. Specifically, this proposed measure reports a SNF's risk-standardized rate of Medicare FFS residents who are discharged to the community following a SNF stay, and do not have an unplanned readmission to an acute care hospital or LTCH in the 31 days following discharge to community, and who remain alive during the 31 days following discharge to community. Community, for this measure, is defined as home/self-care, with or without home health services, based on Patient Discharge Status Codes 01, 06, 81, and 86 on the Medicare FFS claim. 1,2

We have developed three discharge to community measures for IRF, SNF, and LTCH settings, respectively. These measures are conceptualized uniformly across the PAC settings, in terms of the definition of the discharge to community outcome, the approach to risk adjustment, and the measure calculation. It is important to note, though, that each measure is specific to the particular PAC setting (i.e., IRF, SNF, or LTCH); we do not pool PAC patients/residents across settings in the measure development and calculation.

## 2.1.2 Purpose/Rationale for the Measure

Discharge to a community setting is an important health care outcome for many patients/residents for whom the overall goals of post-acute care include optimizing functional improvement, returning to a previous level of independence, and avoiding institutionalization. Returning to the community is also an important outcome for many patients/residents who are not expected to make functional improvement during their PAC stay, and for patients/residents who may be expected to decline functionally due to their medical condition. The discharge to community outcome offers a multi-dimensional view of preparation for community life,

<sup>&</sup>lt;sup>1</sup> Further description of patient discharge status codes can be found, for example, at the following Web page: https://med.noridianmedicare.com/web/jea/topics/claim-submission/patient-status-codes.

<sup>&</sup>lt;sup>2</sup> This definition is not intended to suggest that board and care homes, assisted living facilities, or other settings included in the definition of "community" for the purpose of this measure are the most integrated setting for any particular individual or group of individuals under the Americans with Disabilities Act (ADA) and Section 504.

including the cognitive, physical, and psychosocial elements involved in a discharge to the community.<sup>3,4</sup>

In addition to being an important outcome from a patient/resident and family perspective, patients/residents discharged to community settings, on average, incur lower costs over the recovery episode, compared with those discharged to institutional settings. <sup>5,6</sup> Given the high costs of care in institutional settings, encouraging PACs to prepare patients/residents for discharge to community, when clinically appropriate, may have cost-saving implications for the Medicare program. <sup>7</sup> Also, providers have found that successful discharge to community was a major driver of their ability to achieve savings, where capitated payments for post-acute care were in place. <sup>8</sup> For patients/residents who require long-term care due to persistent disability, discharge to community could result in lower long-term care costs for Medicaid and for patients' or residents' out-of-pocket expenditures. <sup>9</sup>

Analyses conducted for the Assistant Secretary for Planning and Evaluation (ASPE) on PAC episodes, using a 5 percent sample of 2006 Medicare claims, revealed that relatively high average, unadjusted Medicare payments are associated with discharge to institutional settings from IRFs, SNFs, LTCHs or HHAs, as compared with payments associated with discharge to community settings. <sup>10</sup> Average, unadjusted Medicare payments associated with discharge to community settings ranged from \$0 to \$4,017 for IRF discharges, \$0 to \$3,544 for SNF discharges, \$0 to \$4,706 for LTCH discharges, and \$0 to \$992 for HHA discharges. In contrast, payments associated with discharge to non-community settings were considerably higher, ranging from \$11,847 to \$25,364 for IRF discharges, \$9,305 to \$29,118 for SNF discharges, \$12,465 to \$18,205 for LTCH discharges, and \$7,981 to \$35,192 for HHA discharges. <sup>11</sup>

Measuring and comparing facility-level discharge to community rates is expected to help differentiate among facilities with varying performance in this important domain, and to help avoid disparities in care across patient/resident groups. Variation in discharge to community

Doran JP, Zabinski SJ. Bundled payment initiatives for Medicare and non-Medicare total joint arthroplasty patients at a community hospital: bundles in the real world. The Journal of arthroplasty. 2015;30(3):353-355.

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<sup>&</sup>lt;sup>3</sup> El-Solh AA, Saltzman SK, Ramadan FH, Naughton BJ. Validity of an artificial neural network in predicting discharge destination from a postacute geriatric rehabilitation unit. Archives of physical medicine and rehabilitation. 2000;81(10):1388-1393.

<sup>&</sup>lt;sup>4</sup> Tanwir S, Montgomery K, Chari V, Nesathurai S. Stroke rehabilitation: availability of a family member as caregiver and discharge destination. *European journal of physical and rehabilitation medicine*. 2014;50(3):355-362.

Dobrez D, Heinemann AW, Deutsch A, Manheim L, Mallinson T. Impact of Medicare's prospective payment system for inpatient rehabilitation facilities on stroke patient outcomes. *American journal of physical medicine & rehabilitation / Association of Academic Physiatrists.* 2010;89(3):198-204.

<sup>&</sup>lt;sup>6</sup> Gage B, Morley M, Spain P, Ingber M. Examining Post Acute Care Relationships in an Integrated Hospital System. Final Report. RTI International;2009.

lbid.

Newcomer RJ, Ko M, Kang T, Harrington C, Hulett D, Bindman AB. Health Care Expenditures After Initiating Long-term Services and Supports in the Community Versus in a Nursing Facility. Med Care. 2016 Jan 12. *Epub ahead of print*.

Gage B, Morley M, Spain P, Ingber M. Examining Post Acute Care Relationships in an Integrated Hospital System. Final Report. RTI International;2009.

<sup>&</sup>quot; Ibid.

rates has been reported within and across post-acute settings; across a variety of facility-level characteristics, such as geographic location (for example, regional location, urban or rural location), ownership (for example, for-profit or nonprofit), and freestanding or hospital-based units; and across patient-level characteristics, such as race and gender. <sup>12,13,14,15,16,17</sup> Discharge to community rates in the IRF setting have been reported to range from about 60 to 80 percent. <sup>18,19,20,21,22,23</sup> Longer-term studies show that rates of discharge to community from IRFs have decreased over time as IRF length of stay has decreased. <sup>24,25</sup> Greater variation in discharge

Reistetter TA, Karmarkar AM, Graham JE, et al. Regional variation in stroke rehabilitation outcomes. *Archives of physical medicine and rehabilitation*. 2014;95(1):29-38.

El-Solh AA, Saltzman SK, Ramadan FH, Naughton BJ. Validity of an artificial neural network in predicting discharge destination from a postacute geriatric rehabilitation unit. Archives of physical medicine and rehabilitation. 2000;81(10):1388-1393.

March 2015 Report to the Congress: Medicare Payment Policy. Medicare Payment Advisory Commission;2015.

Bhandari VK, Kushel M, Price L, Schillinger D. Racial disparities in outcomes of inpatient stroke rehabilitation. *Archives of physical medicine and rehabilitation*. 2005;86(11):2081-2086.

Chang PF, Ostir GV, Kuo YF, Granger CV, Ottenbacher KJ. Ethnic differences in discharge destination among older patients with traumatic brain injury. Archives of physical medicine and rehabilitation. 2008;89(2):231-236.

Berges IM, Kuo YF, Ostir GV, Granger CV, Graham JE, Ottenbacher KJ. Gender and ethnic differences in rehabilitation outcomes after hip-replacement surgery. *American journal of physical medicine & rehabilitation / Association of Academic Physiatrists.* 2008;87(7):567-572.

Galloway RV, Granger CV, Karmarkar AM, et al. The Uniform Data System for Medical Rehabilitation: report of patients with debility discharged from inpatient rehabilitation programs in 2000-2010. *American journal of physical medicine & rehabilitation / Association of Academic Physiatrists.* 2013;92(1):14-27.

Morley MA, Coots LA, Forgues AL, Gage BJ. Inpatient rehabilitation utilization for Medicare beneficiaries with multiple sclerosis. *Archives of physical medicine and rehabilitation*. 2012;93(8):1377-1383.

Reistetter TA, Graham JE, Deutsch A, Granger CV, Markello S, Ottenbacher KJ. Utility of functional status for classifying community versus institutional discharges after inpatient rehabilitation for stroke. *Archives of physical medicine and rehabilitation*. 2010;91(3):345-350.

Gagnon D, Nadeau S, Tam V. Clinical and administrative outcomes during publicly-funded inpatient stroke rehabilitation based on a case-mix group classification model. *Journal of rehabilitation medicine*. 2005;37(1):45-52.

DaVanzo J, El-Gamil A, Li J, Shimer M, Manolov N, Dobson A. Assessment of patient outcomes of rehabilitative care provided in inpatient rehabilitation facilities (IRFs) and after discharge. Vienna, VA: Dobson DaVanzo & Associates, LLC:2014.

Kushner DS, Peters KM, Johnson-Greene D. Evaluating Siebens Domain Management Model for Inpatient Rehabilitation to Increase Functional Independence and Discharge Rate to Home in Geriatric Patients. *Archives of physical medicine and rehabilitation*. 2015;96(7):1310-1318.

Galloway RV, Granger CV, Karmarkar AM, et al. The Uniform Data System for Medical Rehabilitation: report of patients with debility discharged from inpatient rehabilitation programs in 2000-2010. *American journal of physical medicine & rehabilitation / Association of Academic Physiatrists.* 2013;92(1):14-27.

Mallinson T, Deutsch A, Bateman J, et al. Comparison of discharge functional status after rehabilitation in skilled nursing, home health, and medical rehabilitation settings for patients after hip fracture repair. *Archives of physical medicine and rehabilitation*. 2014;95(2):209-217.

to community rates is seen in the SNF setting, with rates ranging from 31 to 65 percent. <sup>26,27,28,29</sup> In the SNF Medicare FFS population, using CY 2013 national claims data, we found that approximately 44 percent of residents were discharged to the community. A multi-center study of 23 LTCHs demonstrated that 28.8 percent of 1,061 patients who were ventilator-dependent on admission were discharged to home. <sup>30</sup> A single-center study found that 31 percent of LTCH hemodialysis patients were discharged to home. <sup>31</sup> One study noted that 64 percent of beneficiaries who were discharged from the home health episode did not use any other acute or post-acute services paid by Medicare in the 30 days after discharge. <sup>32</sup> However, significant numbers of patients were admitted to hospitals (29 percent) and lesser numbers to skilled nursing facilities (7.6 percent), inpatient rehabilitation facilities (1.5 percent), home health (7.2 percent) or hospice (3.3 percent). <sup>33</sup>

Discharge to community is an actionable health care outcome, as targeted interventions have been shown to successfully increase discharge to community rates in a variety of post-acute settings. 34,35,36,37 Many of these interventions involve discharge planning or specific rehabilitation strategies, such as addressing discharge barriers and improving medical and

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El-Solh AA, Saltzman SK, Ramadan FH, Naughton BJ. Validity of an artificial neural network in predicting discharge destination from a postacute geriatric rehabilitation unit. *Archives of physical medicine and rehabilitation*. 2000;81(10):1388-1393.

Hall RK, Toles M, Massing M, et al. Utilization of acute care among patients with ESRD discharged home from skilled nursing facilities. *Clinical journal of the American Society of Nephrology : CJASN.* 2015;10(3):428-434.

Stearns SC, Dalton K, Holmes GM, Seagrave SM. Using propensity stratification to compare patient outcomes in hospital-based versus freestanding skilled-nursing facilities. *Medical care research and review : MCRR*. 2006;63(5):599-622.

Wodchis WP, Teare GF, Naglie G, et al. Skilled nursing facility rehabilitation and discharge to home after stroke. *Archives of physical medicine and rehabilitation*. 2005;86(3):442-448.

Scheinhorn DJ, Hassenpflug MS, Votto JJ, et al. Post-ICU mechanical ventilation at 23 long-term care hospitals: a multicenter outcomes study. *Chest.* 2007;131(1):85-93.

Thakar CV, Quate-Operacz M, Leonard AC, Eckman MH. Outcomes of hemodialysis patients in a long-term care hospital setting: a single-center study. *American journal of kidney diseases: the official journal of the National Kidney Foundation*. 2010;55(2):300-306.

Wolff JL, Meadow A, Weiss CO, Boyd CM, Leff B. Medicare home health patients' transitions through acute and post-acute care settings. *Medical care*. 2008;46(11):1188-1193.

<sup>&</sup>quot; Ibid.

Kushner DS, Peters KM, Johnson-Greene D. Evaluating Siebens Domain Management Model for Inpatient Rehabilitation to Increase Functional Independence and Discharge Rate to Home in Geriatric Patients. Archives of physical medicine and rehabilitation. 2015;96(7):1310-1318.

Wodchis WP, Teare GF, Naglie G, et al. Skilled nursing facility rehabilitation and discharge to home after stroke. *Archives of physical medicine and rehabilitation*. 2005;86(3):442-448.

Berkowitz RE, Jones RN, Rieder R, et al. Improving disposition outcomes for patients in a geriatric skilled nursing facility. *Journal of the American Geriatrics Society*. 2011;59(6):1130-1136.

Kushner DS, Peters KM, Johnson-Greene D. Evaluating use of the Siebens Domain Management Model during inpatient rehabilitation to increase functional independence and discharge rate to home in stroke patients. *PM & R : the journal of injury, function, and rehabilitation.* 2015;7(4):354-364.

functional status. <sup>38,39,40,41</sup> The effectiveness of these interventions suggests that improvement in discharge to community rates among post-acute care patients/residents is possible through modifying provider-led processes and interventions.

#### 2.1.3 Denominator

The denominator for the discharge to community measure is the risk-adjusted expected number of discharges to community. This estimate includes risk adjustment for patient/resident characteristics with the facility effect removed. The "expected" number of discharges to community is the predicted number of risk-adjusted discharges to community if the same patients/residents were treated at the average facility appropriate to the measure.

The regression model used to calculate the denominator is developed using all non-excluded facility stays in the national data. The denominator is computed in the same way as the numerator, but the facility effect is set at the average. The descriptions of the discharge to community outcome, patient/resident stays included in the measure, and numerator calculation are provided below.

#### 2.1.4 Numerator

The measure does not have a simple form for the numerator and denominator—that is, the risk adjustment method does not make the *observed* number of community discharges the numerator, and a *predicted* number the denominator. The measure numerator is the *risk-adjusted estimate* of the number of patients/residents who are discharged to the community, do not have an unplanned readmission to an acute care hospital or LTCH in the 31-day post-discharge observation window, and who remain alive during the post-discharge observation window. This estimate starts with the observed discharges to community, and is risk adjusted for patient/resident characteristics and a statistical estimate of the facility effect beyond case mix.

The numerator uses a model estimated on full national data specific to the post-acute setting; it is applied to the facility's patient/resident stays included in the measure, and includes the estimated effect of that facility. The prediction equation is based on a logistic statistical model with a two-level hierarchical structure. The patient/resident stays in the model have an indicator of the facility they are discharged from; the effect of the facility is measured as a positive or negative shift in the intercept term of the equation. The facility effects are modeled as belonging to a normal (Gaussian) distribution centered at 0, and are estimated along with the effects of patient/resident characteristics in the model. Numerator details are provided below.

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Kushner DS, Peters KM, Johnson-Greene D. Evaluating Siebens Domain Management Model for Inpatient Rehabilitation to Increase Functional Independence and Discharge Rate to Home in Geriatric Patients. *Archives of physical medicine and rehabilitation*. 2015;96(7):1310-1318.

Wodchis WP, Teare GF, Naglie G, et al. Skilled nursing facility rehabilitation and discharge to home after stroke. *Archives of physical medicine and rehabilitation*. 2005;86(3):442-448.

Berkowitz RE, Jones RN, Rieder R, et al. Improving disposition outcomes for patients in a geriatric skilled nursing facility. *Journal of the American Geriatrics Society*. 2011;59(6):1130-1136.

Kushner DS, Peters KM, Johnson-Greene D. Evaluating use of the Siebens Domain Management Model during inpatient rehabilitation to increase functional independence and discharge rate to home in stroke patients. *PM & R: the journal of injury, function, and rehabilitation.* 2015;7(4):354-364.

Numerator Details: Discharge to Community

Discharge to community is determined based on the "Patient Discharge Status Code" from the PAC claim. Discharge to community is defined as discharge to home/self-care with or without home health services. <sup>42</sup> Table 1 below lists the Patient Discharge Status Codes used to define community.

Table 1
Patient Discharge Status Codes Used to Determine Discharge to Community

Discharge Status Codes Indicating Community Discharge					
01	Discharged to home/self-care (routine discharge)				
06	Discharged/transferred to home under care of organized home health service organization				
81	Discharged to home or self-care with a planned acute care hospital readmission				
86	Discharged/transferred to home under care of organized home health service organization with a planned acute care hospital inpatient readmission				

Numerator Details: Unplanned Readmissions in the 31-Day Post-Discharge Observation Window

A patient/resident who is discharged to the community is considered to have an unfavorable outcome if they have a subsequent unplanned readmission to an acute care hospital or LTCH in the post-discharge observation window, which includes the day of discharge and the 31 days following day of discharge. We identify unplanned readmissions based on the planned readmissions algorithm used in the following post-acute care readmission measures, endorsed by the National Quality Forum (NQF): (i) NQF #2510: Skilled Nursing Facility 30-Day All-Cause Readmission Measure (SNFRM); (ii) NQF #2502: All-Cause Unplanned Readmission Measure for 30 Days Post Discharge from Inpatient Rehabilitation Facilities; (iii) NQF #2512: All-Cause Unplanned Readmission Measure for 30 Days Post Discharge from Long Term Care Hospitals; and (iv) NQF #2380: Rehospitalization During the First 30 Days of Home Health. 43,44,45,46 These readmission measures are based on the Hospital-Wide All-Cause Readmission Measure

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<sup>&</sup>lt;sup>42</sup> Further description of patient discharge status codes can be found, for example, at the following Web page: <a href="https://med.noridianmedicare.com/web/jea/topics/claim-submission/patient-status-codes">https://med.noridianmedicare.com/web/jea/topics/claim-submission/patient-status-codes</a>.

<sup>43</sup> NQF #2510: Skilled Nursing Facility 30-Day All-Cause Readmission Measure (SNFRM). www.qualityforum.org/QPS/2510

<sup>44</sup> NQF #2502: All-Cause Unplanned Readmission Measure for 30 Days Post Discharge from Inpatient Rehabilitation Facilities. www.qualityforum.org/QPS/2502

<sup>45</sup> NQF #2512: All-Cause Unplanned Readmission Measure for 30 Days Post Discharge from Long Term Care Hospitals. www.qualityforum.org/QPS/2512

<sup>46</sup> NQF #2380: Rehospitalization During the First 30 Days of Home Health www.qualityforum.org/QPS/2380

(HWR) (CMS/Yale) (NQF #1789),<sup>47</sup> with some additions made for post-acute settings. The planned readmission definition is based on the claim from the readmission having a code for a procedure that is frequently planned; however, if a principal diagnosis in a specified list of acute diagnoses is present, the readmission is reclassified as unplanned. Readmissions to psychiatric hospitals or units are classified as planned readmissions.

Please note that this measure has been developed with ICD-9 procedure and diagnosis codes. The measure will be revised using the ICD-9 to ICD-10 cross-walk.

Numerator Details: Death in the 31-Day Post-Discharge Observation Window

Patients/residents who are discharged to the community are also considered to have an unfavorable outcome if they die in the post-discharge window, which includes the day of discharge and the 31 days following day of discharge. Death in the post-discharge window is identified based on date of death from Medicare eligibility files.

# 2.1.5 Target Population and Measure Exclusions

The target population for the measure is the group of Medicare FFS patients/residents who are not excluded for the reasons listed below.

#### Measure Exclusions

Exclusions for the discharge to community measure are listed below, along with the rationale for each exclusion. The measure exclusion criteria are determined by processing Medicare claims and eligibility data to determine whether the individual exclusion criteria are met. All measure exclusion criteria are based on administrative data. Only PAC stays that are preceded by a short-term acute care stay in the 30 days prior to the PAC admission date are included in the measure. Stays ending in transfers to the same level of care are excluded.

## 1) Age under 18 years

#### Rationale:

- a. There is limited literature on discharge destination outcomes in this age group;
- b. Patients/residents in this age group represent a different cohort, likely living with their parents, and may be expected to have higher discharge to community rates compared with the rest of the Medicare population; and
- c. Patients/residents in this age group represent a small proportion of the post-acute Medicare FFS population.
- 2) No short-term acute care stay within the 30 days preceding an IRF, SNF, or LTCH admission

<sup>&</sup>lt;sup>47</sup> NQF #1789: Hospital-Wide All-Cause Readmission Measure (HWR) (CMS/Yale). www.qualityforum.org/QPS/1789

Rationale: Acute care claims from the 30 days prior to IRF, SNF, or LTCH admission provide the principal diagnosis and other important patient/resident data for risk adjustment. In IRF, SNF, and LTCH settings, patients/residents without a short-term acute care discharge within the 30 days prior to PAC admission will be excluded from the measure, because important risk adjustment data will be missing.

# 3) Discharges to psychiatric hospital

Rationale: Patients/residents discharged to psychiatric hospital are excluded from the measure because community living at the time of discharge may be potentially inappropriate or unsafe for them due to their mental health or psychiatric condition.

# 4) Discharges against medical advice

Rationale: Patients/residents who discharge themselves against medical advice are excluded because their care plan may not have been fully implemented, and the discharge destination may not reflect the facility's discharge recommendation. Additionally, patients/residents discharged against medical advice may potentially be at higher risk of post-discharge readmissions or death, depending on their medical condition, or due to potential non-adherence or non-compliance with care recommendations.

# 5) Discharges to disaster alternative care sites or federal hospitals

*Rationale:* Patients/residents discharged to disaster alternative care sites are excluded because these discharges are likely influenced by external emergency conditions, and may not represent discretionary discharges by the PAC provider. Discharges to federal hospitals are also excluded.

## 6) Discharges to court/law enforcement

*Rationale:* Patients/residents who are discharged to court or law enforcement are likely ineligible for discharge to the community due to legal restrictions.

# 7) Patients/residents discharged to hospice

#### Rationale:

- a. Patients/residents discharged to hospice care are terminally ill, and have very different goals of care compared with non-hospice patients/residents. For non-hospice patients/residents, the primary goal of post-acute care is to return to baseline, independent living in the community; death is an undesirable outcome in the non-hospice population. For patients/residents discharged to hospice, the goal is to provide them the opportunity to die comfortably, at home or in a facility.
- b. A large proportion of patients/residents discharged to hospice care die in the 31-day window following discharge from the post-acute setting.

- c. The hospice agency, not the post-acute care setting, makes the final decision of discharge to hospice-home or hospice-facility.
- 8) Patients/residents not continuously enrolled in Part A FFS Medicare for the 12 months prior to the post-acute admission date, and at least 31 days after post-acute discharge date

Rationale: Patients/residents not continuously enrolled in Part A FFS Medicare for the 12 months prior to the PAC admission date are excluded because risk adjustment for certain comorbidities requires information on acute inpatient bills for one year prior to post-acute admission. Patients/residents not continuously enrolled in Part A FFS Medicare for at least 31 days after post-acute discharge are excluded because readmissions and death must be observable in the 31-day post-discharge period. Patients/residents without Part A coverage or those who are enrolled in Medicare Advantage plans will not have complete inpatient claims in the system.

9) Patients/residents whose prior short-term acute care stay was for non-surgical treatment of cancer

*Rationale:* Patients/residents whose prior short-term acute care stay was for non-surgical treatment of cancer are excluded because they have a different trajectory for recovery after discharge, with a high mortality rate.<sup>48</sup> Exclusion of these patients/residents is consistent with the hospital-wide and post-acute readmission measures.

10) Post-acute stays that end in transfer to the same level of care

*Rationale:* Post-acute stays that end in transfer to the same level of care are excluded from the measure because their post-acute episode has not ended. For a post-acute episode that involves transfer to the same level of care, only the final post-acute provider is included in the measure.

11) Post-acute stays with claims data that are problematic (e.g., anomalous records for stays that overlap wholly or in part, or are otherwise erroneous or contradictory)

*Rationale:* This measure requires accurate information from the post-acute stay and prior short-term acute care stay in the elements used for risk adjustment. No-pay post-acute stays involving exhaustion of Part A benefits are also excluded.

12) Planned discharges to an acute or LTCH setting

*Rationale:* For the IRF and SNF settings, planned discharges to an acute care hospital or LTCH will be excluded. For the LTCH setting, planned discharges to an

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<sup>48</sup> NQF #1789: Hospital-Wide All-Cause Readmission Measure (HWR) (CMS/Yale). www.qualityforum.org/QPS/1789

acute care hospital will be excluded. (Note that, in the LTCH setting, transfer to another LTCH is excluded because it represents a transfer to the same level of care).

13) Medicare Part A benefits exhausted

*Rationale:* Patients/residents who have exhausted their Medicare Part A coverage during the PAC stay are excluded because the discharge destination decision may be related to exhaustion of benefits.

14) Patients/residents who received care from a facility located outside of the United States, Puerto Rico or a U.S. territory

Rationale: Patients/residents who received care from foreign facilities may not have complete inpatient claims in the system, and these facilities may not be subject to policy decisions related to this quality measure.

15) Swing Bed Stays in Critical Access Hospitals (SNF setting only)

Rationale: Critical access hospital (CAH) swing bed stays are excluded from the SNF setting measure. This is because CAH swing bed facilities are not required to submit quality data under the SNF QRP, and are exempt from the SNF Prospective Payment System (PPS). Note that non-CAH swing bed stays are included in the measure, because non-CAH swing bed facilities are required to submit quality data under the SNF QRP and are subject to the SNF PPS.

#### 2.1.6 Data Sources

This measure is based on Medicare FFS administrative claims, and uses data in the Medicare eligibility files and inpatient claims. The eligibility files provide information such as date of birth, date of death, sex, reasons for Medicare eligibility, periods of Part A coverage, and periods in the Medicare FFS program. The data elements from the Medicare FFS claims are those basic to the operation of the Medicare payment systems and include data such as date of admission, date of discharge, diagnoses, procedures, indicators for use of dialysis services, and indicators of whether the Part A benefit was exhausted. The inpatient claims data files contain patient/resident-level PAC and other hospital records. No data beyond the bills submitted in the normal course of business are required from providers for the calculation of this measure.

# IRF & LTCH Measure Data Sources

The following are the specific files used for the IRF and LTCH measures and links to their documentation:

Medicare Inpatient claims (Standard Analytical Files), Index PAC claims

Documentation for the Medicare claims data is provided online by ResDAC. The following web page includes data dictionaries for the Standard analytical files (Inpatient RIF): http://www.resdac.org/cms-data/files/ip-rif/data-documentation

Medicare Enrollment Database - Information about the Enrollment Database may be found at: <a href="http://aspe.hhs.gov/datacncl/datadir/cms.htm">http://aspe.hhs.gov/datacncl/datadir/cms.htm</a>

Medicare Denominator files - Documentation available at: <a href="http://www.cms.gov/Research-Statistics-Data-and-Systems/Files-for-Order/IdentifiableDataFiles/DenominatorFile.html">http://www.cms.gov/Research-Statistics-Data-and-Systems/Files-for-Order/IdentifiableDataFiles/DenominatorFile.html</a>

#### SNF Measure Data Sources

The following are the specific files used for the SNF measure and links to their documentation:

Medicare Inpatient Claims (MedPAR RIF), Index SNF Claims:

Documentation for the Medicare claims data is provided online by the CMS contractor, Research Data Assistance Center (ResDAC) at the University of Minnesota. The following web page includes data dictionaries for the MedPAR RIF: <a href="http://www.resdac.org/cms-data/files/medpar-rif">http://www.resdac.org/cms-data/files/medpar-rif</a>

Medicare Enrollment Database - Information about the Enrollment Database may be found at: <a href="http://aspe.hhs.gov/datacncl/datadir/cms.htm">http://aspe.hhs.gov/datacncl/datadir/cms.htm</a>

Medicare Denominator files - Documentation available at: <a href="http://www.cms.gov/Research-Statistics-Data-and-Systems/Files-for-Order/IdentifiableDataFiles/DenominatorFile.html">http://www.cms.gov/Research-Statistics-Data-and-Systems/Files-for-Order/IdentifiableDataFiles/DenominatorFile.html</a>

## 2.1.7 Measure Time Window

Time windows vary across settings due to variation in facility sizes across settings. The measure time window is two years in IRF and LTCH settings, and one year in the SNF setting. Specific measure time window descriptions for each setting are provided below.

*IRF and LTCH Time Windows:* In the IRF and LTCH settings, the measure is calculated using two years of data. All IRF and LTCH stays during the two-year time window, except those that meet the exclusion criteria, are included in the measure. For patients with multiple stays during the two-year time window, each stay is eligible for inclusion in the measure. Data from CY 2012-2013 were used to develop this measure.

*SNF Time Window:* In the SNF setting, the measure is calculated using one year of data. All SNF stays during the one-year time window, except those that meet the exclusion criteria, are included in the measure. For SNF residents with multiple SNF stays during the one year window, each stay is eligible for inclusion in the measure. Data from CY 2013 were used to develop this measure.

# 2.1.8 Statistical Risk Model and Risk Adjustment Covariates

We used a hierarchical logistic regression method to predict the probability of discharge to community. Patient/resident characteristics related to discharge and a marker for the specific discharging facility are included in the equation. The equation is hierarchical in that both

individual patient/resident characteristics are accounted for, as well as the clustering of patient/resident characteristics by facility. The statistical model estimates both the average predictive effect of the patient/resident characteristics across all facilities, and the degree to which each facility has an effect on discharge to community that differs from that of the average facility. The facility effects are assumed to be randomly distributed around the average (according to a normal distribution). When computing the facility effect, hierarchical modeling accounts for the known predictors of discharge to community, on average, such as patient/resident characteristics, the observed facility rate, and the number of facility stays eligible for inclusion in the measure. The estimated facility effect is determined mostly by the facility's own data if the number of patient/resident discharges is relatively large (as the estimate would be relatively precise), but is adjusted toward the average if the number of patient/resident discharges is small (as that would yield a less precise estimate).

## We used the following model:

Let  $Y_{ij}$ , denote the outcome (equal to 1 if patient/resident i is discharged to community, 0 otherwise) for a patient/resident i at facility j;  $Z_{ij}$  denotes a set of risk adjustment variables. We assume the outcome is related to the risk adjusters via a logit function with dispersion:

$$\begin{split} logit(Prob(Y_{ij} = 1)) &= \alpha_j + \beta*Z_{ij} + \ \epsilon_{ij} \\ \alpha_j &= \mu + \omega_j \ , \ \omega_j \sim N(0, \tau^2) \end{split} \eqno(1)$$

where  $Z_{ij} = (Z_1, Z_2, ... Z_k)$  is a set of k patient/resident-level risk adjustment variables;  $\alpha_j$  represents the facility-specific intercept;  $\mu$  is the adjusted average outcome across all facilities;  $\tau^2$  is the between-facility variance component; and  $\epsilon \sim N(0, \sigma^2)$  is the error term. The hierarchical logistic regression model is estimated using SAS software (PROC GLIMMIX: SAS/STAT User's Guide, SAS Institute Inc.).

The estimated equation is used twice in the measure. The sum of the probabilities of discharge to community of all patients/residents in the facility measure, including both the effects of patient/resident characteristics and the facility, is the "predicted number" of discharges to community after adjusting for the facility's case mix. The same equation is used without the facility effect to compute the "expected number" of discharges to community for the same patients/residents at the average facility. The ratio of the predicted-to-expected number of discharges to community is a measure of the degree to which discharges to community are higher or lower than what would otherwise be expected. This standardized risk ratio is then multiplied by the mean discharge to community rate for all facility stays for the measure, yielding the risk-standardized discharge to community rate for each facility. Please note that the estimation procedure is recalculated for each measurement period. Re-estimating the models for each measurement period allows the estimated effects of the patient/resident characteristics to vary over time as patient/resident case-mix and medical treatment patterns change.

Risk adjustment variables include demographic and eligibility characteristics; principal diagnoses; types of surgery or procedures from the prior short-term acute care stay; comorbidities; length of stay and intensive care utilization from the prior short-term acute care stay; dialysis in the prior acute stay; and number of prior hospitalizations in the year preceding

the PAC admission. Risk adjustment variable descriptions are provided below. See Appendix Table 1 for the full list of variables in the risk adjustment models.

- 1) Age and sex groups.
- 2) End stage renal disease (ESRD) or disability as original reason for entitlement.
- 3) Principal diagnosis (Clinical Classifications Software (CCS) groups) from the prior acute stay in the past 30 days. The ICD-9 codes from the prior acute claim are grouped clinically using the CCS for ICD-9 diagnoses developed by the Agency for Healthcare Research and Quality (AHRQ).<sup>49</sup>
- 4) Case-Mix Groups (in the IRF model).
- 5) Surgical procedure categories (if present) based on the prior acute stay in the past 30 days. The procedures are grouped using the CCS classes for ICD-9 procedures developed by AHRQ.
- 6) Dialysis in prior acute stay where ESRD not indicated.
- 7) Indicator for ESRD status.
- 8) Length of prior acute hospital stay in days, for patients/residents whose prior acute stay was in a non-psychiatric hospital (categorical variables are used to account for nonlinearity); indicator of prior psychiatric hospital stay for patients/residents whose prior acute stay was in a psychiatric hospital.
- 9) Number of intensive/cardiac care days during the prior acute stay (in the LTCH model).
- 10) Ventilator use during the post-acute stay (in the LTCH and SNF models).
- 11) Comorbidities (Hierarchical Condition Categories) (based on prior acute stay in the past 30 days or based on a one year look back, depending on the specific comorbidity). Comorbidities are clustered using the Hierarchical Condition Categories [HCC] groups used by CMS.<sup>50</sup>
- 12) Number of prior acute hospital discharges in the past year, not including the hospitalization in the 30 days prior to the post-acute stay.

50 CMS-HCC Mappings of ICD-9 Codes: Mappings are included in the software at the following website: http://www.cms.gov/Medicare/Health-Plans/MedicareAdvtgSpecRateStats/Risk-Adjustors.html

<sup>49</sup> AHRQ CCS groupings of ICD-9 codes - Documentation available at: <a href="http://www.hcup-us.ahrq.gov/toolssoftware/ccs/ccs.jsp">http://www.hcup-us.ahrq.gov/toolssoftware/ccs/ccs.jsp</a>

us.ahrq.gov/toolssoftware/ccs/ccs.jsp

# 2.1.9 Measure Calculation Algorithm

The following steps describe the calculation algorithm/measure logic for the discharge to community measures:

- Step 1: Identify patients/residents meeting the criteria for the target population, after applying measure exclusions.
- Step 2: Identify patients/residents meeting the numerator criteria, i.e., discharge to community, no unplanned readmissions on the day of discharge or in the 31 days following discharge, and no death on the day of discharge or in the 31 days following discharge.
- Step 3: Identify presence or absence of risk adjustment variables for each patient/resident.
- Step 4: Calculate the predicted and expected number of discharges to community for each facility using the hierarchical logistic regression model.

The predicted number of discharges to community for each facility is calculated as the sum of the predicted probability of discharge to community for each patient/resident discharged from the facility and included in the measure, including the facility-specific effect.

To calculate the predicted number of discharges to community, pred<sub>j</sub>, for index facility stays at facility<sub>j</sub>, we used the following equation:

$$pred_{i} = \sum logit^{-1}(\mu + \omega_{i} + \beta * Z_{ii})$$
 (2)

where the sum is over all stays in facility<sub>i</sub>, and  $\omega_i$  is the random intercept.

To calculate the expected number exp<sub>i</sub>, we used the following equation:

$$exp_j = \Sigma logit^{-1} (\mu + \beta * Z_{ij})$$
 (3)

Step 5: Calculate the standardized risk ratio for each facility, as the ratio of the predicted to expected number of discharges to community.

To calculate the facility-wide standardized risk ratio,  $SRR_j$ , we used the following equation:

$$SRR_i = pred_i / exp_i \tag{4}$$

Step 6: Calculate the risk-standardized discharge to community rate for each facility.

To aid interpretation, the facility-wide standardized risk ratio, SRR<sub>j</sub>, obtained from equation (4) is then multiplied by the overall national raw

discharge to community rate for all facility stays,  $\bar{Y}$ , to produce the facility-wide risk-standardized discharge to community rate (RSR<sub>j</sub>).

To calculate the risk-standardized discharge to community rate for each facility, we used the following equation:

$$RSR_{i} = SRR_{i} * \bar{Y}$$
 (5)

NOTE: Because the statistic described in Step six is a complex function of parameter estimates, re-sampling and simulation techniques (e.g., bootstrapping) may be necessary to derive a confidence interval estimate for the final risk-standardized rate, to characterize the uncertainty of the estimate.

See **Appendix 1, Table 1-1** for risk adjustment model results. Distribution of facility-level discharge to community rates is provided in **Appendix 1, Table 1-2** and **Appendix 1, Figure 1-1**.

# 2.2 Potentially Preventable 30-Day Post-Discharge Readmission Measure for Skilled Nursing Facility (SNF) Quality Reporting Program (QRP)

# 2.2.1 Measure Description

This set of potentially preventable readmission (PPR) measures for post-acute care (PAC) estimates the risk-standardized rate of unplanned, potentially preventable readmissions for patients/residents (Medicare fee-for-service [FFS] beneficiaries) who receive services in one of the following post-acute care provider types: skilled nursing facilities (SNFs), inpatient rehabilitation facilities (IRFs), and long-term care hospitals (LTCH). This measure is conceptualized uniformly across the PAC settings, in terms of the definition of the PPR outcome, the approach to risk adjustment, and the measure calculation.

These outcome measures reflect readmission rates for patients/residents who are readmitted to a short-stay acute-care hospital or an LTCH with a principal diagnosis considered to be unplanned and potentially preventable.

Four PPR PAC measures are being developed.

- 1) Potentially Preventable 30-Day Post-Discharge Readmission Measure for Skilled Nursing Facility Quality Reporting Program (IMPACT)
- 2) Potentially Preventable 30-Day Post-Discharge Readmission Measure for Inpatient Rehabilitation Facility Quality Reporting Program (IMPACT)
- 3) Potentially Preventable 30-Day Post-Discharge Readmission Measure for Long-Term Care Hospital Quality Reporting Program (IMPACT)
- 4) Potentially Preventable Within Stay Readmission Measure for Inpatient Rehabilitation Facilities. Note: This measure is not being developed to meet IMPACT Act requirements.

Three of these measures assess PPR within a 30-day window following discharge from PAC—one measure for each PAC setting (i.e. SNF, IRF, and LTCH)—and are being developed to meet the requirements of the IMPACT Act. An additional IRF measure assesses PPR during the IRF stay (referred to as the within-stay window), and is being developed for use in the IRF Quality Reporting Program. There is also a related SNF measure of potentially preventable hospital readmissions, which is being developed along with the measures cited above in order to meet the requirements of the Protecting Access to Medicare Act of 2014; this measure is detailed in a separate document.

Each measure calculates a risk-adjusted PPR rate for each PAC provider. This is derived by first calculating a standardized risk ratio -- the predicted number of readmissions at the PAC provider (facility) divided by the expected number of readmissions for the same patients/residents if treated at the average PAC provider. The standardized risk ratio is then multiplied by the mean readmission rate in the population (i.e., all Medicare FFS patients/residents included in the measure) to generate the PAC provider-level standardized readmission rate of potentially preventable readmissions.

For these PPR measures, readmissions that are usually for planned procedures are not counted as being potentially preventable (see details below).

## 2.2.2 Purpose/Rationale for the Measure

Hospital readmissions among the Medicare population are common, costly, and often preventable. <sup>51,52</sup> The Medicare Payment Advisory Commission (MedPAC) and a study by Jencks et al. estimated that 17-20 percent of Medicare beneficiaries discharged from the hospital were readmitted within 30 days. Among these hospital readmissions, MedPAC has estimated that 76 percent were considered potentially avoidable--associated with \$12 billion in Medicare expenditures. <sup>53,54</sup>

The Centers for Medicare & Medicaid Services (CMS) has addressed the high rates of hospital readmissions for the acute care hospital setting and more recently, among post-acute care providers. For example, CMS developed the following all-cause readmission measures: All-Cause Unplanned Readmission Measure for 30 days Post Discharge from Inpatient Rehabilitation Facilities (IRFs), All-Cause Unplanned Readmission Measure for 30 days Post Discharge from Long-Term Care Hospitals (LTCHs), and the Skilled Nursing Facility (SNF) 30-Day All-Cause Readmission Measure (NQF #2380, #2502, #2512, and #2510, respectively). These measures were endorsed by the National Quality Forum (NQF). The IRF and LTCH measures were adopted for their respective quality reporting programs for public reporting, and

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<sup>&</sup>lt;sup>51</sup> Friedman, B. and J. Basu, The rate and cost of hospital readmissions for preventable conditions. Med Care Res Rev, 2004. **61**(2): p. 225-40.

<sup>&</sup>lt;sup>52</sup> Jencks, S.F., M.V. Williams, and E.A. Coleman, Rehospitalizations among Patients in the Medicare Fee-for-Service Program. New England Journal of Medicine, 2009. **360**(14): p. 1418-1428.

<sup>&</sup>lt;sup>53</sup> *Ibid*.

MedPAC, Payment policy for inpatient readmissions, in Report to the Congress: Promoting Greater Efficiency in Medicare. 2007: Washington D.C. p. 103-120.

<sup>55</sup> National Quality Forum., All-Cause Admissions and Readmissions Measures. April 2015. p. 1-319.

the SNF measure was adopted for value-based purchasing. The NQF-endorsed measures focus on all-cause readmissions and are not cross-setting in that the specifications differ by measure.

Current work is focused on the development of potentially preventable hospital readmission measures for post-acute care, as directed by Congress through the *Improving Medicare Post-Acute Care Transformation Act of 2014* (IMPACT Act). The IMPACT Act requires the development and submission of standardized data from post-acute care settings with the intent for cross-setting quality comparison to promote patient-centeredness. <sup>56</sup> This includes the requirement to develop and implement measures to reflect all-condition risk-adjusted potentially preventable hospital readmission rates.

#### 2.2.3 Denominator

The denominator for the PPR measures is computed the same way as the numerator, but the facility effect is set at the average. The details of the readmission types counted in the numerator and the patients/residents who are included in the measures are below.

For the eligible PAC stays at each facility, the measure denominator is the risk-adjusted expected number of readmissions. This estimate includes risk adjustment for patient/resident characteristics with the facility effect removed. The "expected" number of readmissions is the predicted number of risk-adjusted readmissions if the same patients/residents were treated at the average PAC provider appropriate to the measure.

This population, like that of the numerator, is the group of Medicare FFS PAC patients/residents who are not excluded for the reasons below.

Denominator Exclusions: SNF, IRF, and LTCH Post-Discharge Measures

The post-PAC discharge PPR measures are based on Medicare FFS claims data and include PAC discharges to non-hospital post-acute levels of care or to the community. The observation window is 30-days after discharge from a PAC facility; this window of observation excludes the day of discharge and the day thereafter (i.e. the 30 days starts 2 days after the discharge date). Stays ending in transfers to the same level of care or acute hospitals are excluded. Only PAC stays where patients/residents had a short-term acute care stay within 30 days prior to the PAC admission date are included in the measures. Prior proximal hospital stays include an inpatient admission to an acute care hospital (including IPPS, CAH, or a psychiatric hospital).

1) Patients/residents who died during the SNF/IRF/LTCH stay.

*Rationale*: The PPR measures are not relevant for patients/residents who died during their PAC stay because there is no post-PAC discharge period to observe.

2) Patients/residents less than 18 years old.

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<sup>&</sup>lt;sup>56</sup> United States Congress., H.R. 4994. *IMPACT Act of 2014*. 2014: United States of America. p. 1-19

*Rationale:* Patients/residents under 18 years old are not included in the target population for this measure. Pediatric patients/residents are relatively few and may have different patterns of care than adults.

3) Patients/residents who were transferred at the end of a stay to another SNF/IRF/LTCH or short-term acute care hospital.

Rationale: SNF, IRF, or LTCH patients/residents who were transferred to another SNF/IRF/LTCH or short-term acute-care hospital are excluded from this measure because the transfer suggests that either their SNF/IRF/LTCH treatment has not been completed or that their condition worsened, requiring a transfer (i.e. readmission) back to the acute care setting. The intent of these measures is to follow patients/residents deemed well enough to be discharged to a less intensive care setting (i.e., discharged to less intense levels of care or to the community).

4) Patients/residents who were not continuously enrolled in Part A FFS Medicare for the 12 months prior to the SNF/IRF/LTCH admission date, and at least 30 days after SNF/IRF/LTCH discharge date.

Rationale: The adjustment for certain comorbid conditions in the measures requires information on acute inpatient claims for one year prior to the SNF/IRF/LTCH admission, and readmissions must be observable in the observation window following discharge. Patients/residents without Part A coverage or who are enrolled in Medicare Advantage plans will not have complete inpatient claims in the system.

5) Patients/residents who did not have a short-term acute-care stay within 30 days prior to a SNF/IRF/LTCH admission date.

*Rationale:* These measures require information from the prior short-term acute-care stay in the elements used for risk adjustment.

6) Patients/residents discharged against medical advice (AMA).

*Rationale:* Patients/residents discharged AMA are excluded because these patients/residents have not completed their full course of treatment in the opinion of the facility.

7) Patients/residents for whom the prior short-term acute-care stay was for nonsurgical treatment of cancer.

Rationale: Consistent with the Hospital Wide Readmission (HWR) Measure (NQF #1789), patients/residents for whom the prior short-term acute-care stay was for nonsurgical treatment of cancer are excluded because these patients/residents were identified as following a very different trajectory after discharge, with a particularly high mortality rate.

8) Patients/residents who were transferred to a federal hospital from the PAC facility.

- *Rationale:* Patients/residents who are transferred to federal hospitals will not have complete inpatient claims in the system.
- 9) Patients/residents who received care from a provider located outside of the United States, Puerto Rico, or a U.S. territory.
  - *Rationale:* Patients/residents who received care from foreign providers may not have complete inpatient claims in the system, and these providers may not be subject to the same policy decisions related to readmissions.
- 10) SNF/IRF/LTCH stays with data that are problematic (e.g., anomalous records for hospital stays that overlap wholly or in part, or are otherwise erroneous or contradictory). This also includes SNF stays for residents who exhausted their Medicare benefits for SNF coverage.
  - Rationale: This measure requires accurate information from the SNF/IRF/LTCH stay and prior short-term acute-care stays, for the elements used in risk adjustment. No-pay PAC stays involving exhaustion of Part A benefits are also excluded.
- 11) SNF stays in which the prior proximal hospitalization was for pregnancy.

Rationale: This is a very atypical reason for beneficiaries to be admitted to SNFs.

#### 2.2.4 Numerator

As described, the index PAC admission must have occurred within up to 30 days of discharge from a prior proximal hospital stay (including IPPS, CAH, or a psychiatric hospital). Hospital readmissions include readmissions to a short-stay acute-care hospital or an LTCH, with a diagnosis considered to be unplanned and potentially preventable. Note: Readmissions to inpatient psychiatric facilities are considered planned and not counted for the purposes of this measure.

The numerators of these measures are mathematically related to the number of patients/residents in the target population who have the event of a potentially preventable, unplanned readmission (PPR definitions and planned readmissions are further described below) during the specific readmission window (i.e. 30-day post-PAC discharge). Each measure includes only one readmission window, as described above.

The measures do not have a simple form for the numerator and denominator—that is, the risk adjustment method does not make the observed number of readmissions the numerator, and a predicted number the denominator. Instead, the numerator is the risk-adjusted estimate of the number of unplanned readmissions that occurred within 30 days of PAC discharge. This estimate starts with the observed readmissions, and is then risk-adjusted for patient/resident characteristics and a statistical estimate of the PAC provider's effect, beyond patient/resident case mix.

The prediction equations are based on a logistic statistical model with a 2-level hierarchical structure. The patient/resident stays in the model have an indicator as to which PAC provider they are discharged from and the effect of the provider is measured as a positive or

negative shift in the intercept term of the equation. The facility effects are modeled as belonging to a normal (Gaussian) distribution centered at 0, and are estimated along with the effects of patient characteristics in the model.

The data are from Medicare FFS inpatient claims, and eligibility and enrollment data. Because this measure is claims-based, there is no additional data collection or submission burden for providers.

See below for more details on the data sources.

NOTE: These measures were developed with ICD-9 procedure and diagnosis codes. ICD-10 was implemented on October 1, 2015; when we calculate this measure using data from calendar year 2015, we will use ICD-10 codes. A preliminary list of the PPR definition using ICD-10 codes can be found in **Appendix 2, Table 2-2**.

#### Numerator Details: Readmissions Counted in Measures

# PPR Definitions

Some general methods and algorithms have been developed to assess potentially avoidable or preventable hospitalizations and readmissions for the general Medicare population, such as the Agency for Healthcare Research and Quality's (AHRQ) Prevention Quality Indicators (PQI), approaches developed by and for MedPAC, and proprietary approaches, such as the 3M<sup>TM</sup> algorithm for Potentially Preventable Readmissions.<sup>57,58,59</sup> However, there is no consensus on how to define potentially avoidable or preventable readmissions, especially among Medicare beneficiaries who utilize PAC services including SNF, IRF, and LTCH. Recent work led by Kramer et al. for MedPAC identified 13 conditions that were deemed potentially preventable among the SNF and IRF populations;<sup>60,61</sup> however, these conditions did not differ by PAC setting or readmission window (i.e. during the PAC stay or post-PAC discharge). To support the development of potentially preventable hospital readmission measures among patients/residents who use PAC, measure development contractors (RTI International and Abt Associates) have developed an approach to define potentially preventable readmissions, building on existing research in this area, and are developing measures to address this high priority area.

<sup>57</sup> Goldfield, N.M., Elizabeth; Hughes, John; Tang, Ana; Eastman, Beth; Rawlins, Lisa; Averill, Richard, Identifying Potentially Preventable Readmissions. Health Care Financing Review, 2008. 30(1): p. 75-91.

<sup>&</sup>lt;sup>58</sup> Agency for Healthcare Research and Quality. *Prevention Quality Indicators Overview*. 2008.

<sup>&</sup>lt;sup>59</sup> MedPAC, Online Appendix C: Medicare Ambulatory Care Indicators for the Elderly, in Report to the Congress: Medicare Payment Policy. 2011. p. 7-11.

<sup>60</sup> Kramer, A.L., Michael; Fish, Ron; Min, Sung-Joon, *Development of Potentially Avoidable Readmission and Functional Outcome SNF Quality Measures*. 2014. p. 1-75.

Kramer, A.L., Michael; Fish, Ron; Min, Sung-joon, Development of Inpatient Rehabilitation Facility Quality Measures: Potentially Avoidable Readmissions, Community Discharge, and Functional Improvement. 2014. p. 1-42.

The literature shows that some hospital readmissions can be prevented, and that many of these readmissions occur in the context of PAC, including SNF, IRF, and LTCH.<sup>62,63</sup> For certain diagnoses, proper care and management of patients' or residents' conditions (in the facility or by primary care following discharge) along with appropriate, clearly explained and implemented discharge instructions and referrals, can often prevent a patient's or resident's readmission to the hospital. Identifying these PPR conditions will assist healthcare providers' efforts to improve quality of care and coordination across the care continuum.

In order to develop PPR definitions for PAC, we conducted a comprehensive environmental scan to identify studies and previously published methodologies related to potentially preventable hospitalizations and hospital readmissions. The evidence specific to PAC is limited, and we found substantial variation across methodologies for defining potentially preventable hospitalizations or readmissions. Based on this scan, we compiled a list of all PPR conditions described in the literature. This list had considerable overlap with the Ambulatory Care Sensitive Conditions (ACSC) / PQI, developed by the AHRQ.

We used the ACSC approach as the starting point for this work. Given clinical evidence that these conditions can be avoided with appropriate access to high quality ambulatory care, we found that a majority of these conditions reflect reasons for readmissions that would be considered potentially preventable.<sup>64</sup> We extended this logic to both the within-PAC stay readmission window and 30-day post-PAC discharge window.

In addition, this PPR definition was informed by empirical analyses. Specifically, we analyzed Medicare claims data to identify the most frequent diagnoses associated with hospital readmissions among patients/residents that received post-acute care. We evaluated whether these common causes for readmission could also be considered potentially preventable, by applying the working conceptual definition for PPR explained above, to each of the diagnoses found in the claims analysis. Some conditions such as pressure ulcers, were not on either the ACSC list or in the preliminary data analyses. However, the literature strongly suggests that readmissions for these conditions can be prevented with close monitoring from healthcare providers and under appropriate ambulatory care.

In developing these sets of PPR conditions, we grouped them based on clinical rationale, as follows:

- 1) Inadequate management of chronic conditions
- 2) Inadequate management of infections

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West, J.R., et al., *Determinants of preventable readmissions in the United States: a systematic review.* Implement Sci, 2010. 5: p. 88.

van Walraven, C., A. Jennings, and A.J. Forster, *A meta-analysis of hospital 30-day avoidable readmission rates.* J Eval Clin Pract, 2012. **18**(6): p. 1211-1218.

<sup>64</sup> AHRQ Quality Indicators—Guide to Prevention Quality Indicators: Hospital Admission for Ambulatory Care Sensitive Conditions. Rockville, MD: Agency for Healthcare Research and Quality, 2001. AHRQ Pub. No. 02-R0203.

- 3) Inadequate management of other unplanned events
- 4) Inadequate injury prevention

We sought technical expert and detailed clinical input on these definitions and overall approach. The Technical Expert Panel's (TEP) consensus was that it is feasible to develop uniform definitions that may be applied to all PAC providers. Based on TEP feedback, we substantially revised the definitions to remove several proposed PPR conditions (for example, we excluded several chronic conditions included in the ACSC approach, such as readmissions for long-term complications of diabetes) and, in some cases, added new PPR conditions based on TEP input, such as influenza. In instances where no clear consensus was reached among TEP members (e.g., urinary tract infection, septicemia) we deferred to clinical expertise from the measure development team along with results from our environmental scan which suggested that these conditions were appropriate to consider as potentially preventable.

**Appendix 2, Table 2-1** summarizes the set of conditions we considered potentially preventable for the 30-day post-PAC discharge readmission window based on TEP input. The list of PPR conditions is organized by the clinical rationale for each condition's inclusion on this list.

In order for a readmission to be considered potentially preventable, it must be coded as the principal diagnosis on the readmission claim. However, there are some exceptions based on the PQI specifications, as noted in the appendices (see dehydration conditions).

#### Planned Readmissions

These measures are focused on readmissions that are potentially preventable and *unplanned*. Thus, planned readmissions are not counted in the numerator—PPRs are only counted in the numerator if the readmission is considered unplanned. Planned readmissions are defined largely by the definition used for the HWR measure, and were revised to include additional procedures determined suitable for PAC, with input from a Technical Expert Panel convened by CMS contractors (RTI International and Abt Associates). Both are described in greater detail below. ICD-9 codes for these additional procedures were identified by a certified coder.

If a readmission claim contains a code for a procedure that is frequently a planned procedure, then that readmission is designated to be a planned readmission. However, the readmission is reclassified as unplanned if the claim also contains a code indicating one or more acute diagnoses from a specified list, which can be found in **Appendix 2, Table 2-6**.

Appendix 2, Table 2-7 presents the list of codes for procedures identified as "planned" for PAC, which were not included in the CMS Planned Readmission Algorithm at the time of its development. These procedures and diagnoses are currently defined by ICD-9 procedure and diagnosis codes grouped by the Clinical Classification Software (CCS), developed by the AHRQ. They are included as full CCS classes where appropriate, or by individual codes, if necessary. Readmissions to psychiatric hospitals or units are also classified as planned readmissions.

The Appendix includes details on the planned readmission definitions, including the CMS Planned Readmission Algorithm version 3.0 (**Appendix 2, Figure 2-1 and Tables 2-3 to 2-6**) and a table summarizing the additional planned readmissions added for PAC (**Appendix 2, Table 2-7**). Note this approach is consistent with that used for the NQF-endorsed SNF, IRF, and LTCH all-cause readmission measures (NQF #2510, 2502, and 2512, respectively).

## Readmission Time Frames

The conceptual definition for PPR hinges on the readmission window timeframe. We considered two readmission windows in this work: 1) within-PAC stay and 2) 30 days post-PAC discharge.

For the within-PAC stay window, potentially preventable readmissions should be avoidable with sufficient medical monitoring and appropriate patient/resident treatment. For patients/residents in the 30-day post-PAC discharge period, a potentially preventable readmission refers to a readmission that should be avoidable with adequately planned, explained, and implemented post discharge instructions, including the establishment of appropriate follow-up ambulatory care.

**Table 2** below summarizes the specific readmission windows that are being developed for each PAC potentially preventable hospital readmission measure. As noted, this reflects the current PPR measures under development; however additional measures may be considered in future work. At this time, CMS is only developing a PPR within stay measure for IRFs. Due to data limitations, specifically that claims are not generated for short program interruptions (<4 days) from LTCHs, CMS is unable to develop a PPR within-stay measure for LTCHs.

Table 2
PAC Readmission Windows for Potentially Preventable Hospital Readmission Measure
Development

PAC	Within stay	30-days post PAC discharge (IMPACT Act Measures)
SNF		X
IRF	X	X
LTCH		X

#### Other Documentation

AHRQ CCS groupings of ICD-9 codes: Documentation available at: <a href="http://www.hcup-us.ahrq.gov/toolssoftware/ccs/ccs.jsp">http://www.hcup-us.ahrq.gov/toolssoftware/ccs/ccs.jsp</a>

CMS-HCC Mappings of ICD-9 Codes: Mappings are included in the software at the following website: <a href="http://www.cms.gov/Medicare/Health-Plans/MedicareAdvtgSpecRateStats/Risk-Adjustors.html">http://www.cms.gov/Medicare/Health-Plans/MedicareAdvtgSpecRateStats/Risk-Adjustors.html</a>

#### 2.2.5 Data Sources

All measures are based on administrative claims data.

SNF Measure Data Sources: This measure is for Medicare beneficiaries and uses the data in the Medicare eligibility files and inpatient claims data. The eligibility files provide information on date of birth, sex, reasons for Medicare eligibility, periods of Part A coverage, and periods in the fee-for-service program. The data elements from the Medicare FFS claims are those basic to the operation of the Medicare payment systems and include date of admission, date of discharge, diagnoses, procedures, indicators for use of dialysis services, and indicators of whether the Part A benefit was exhausted. The inpatient claims data files contain beneficiary-level SNF and other hospital records. No data beyond the bills submitted in the normal course of business are required from providers for the calculation of this measure. The following are the specific files and links to their documentation:

Medicare Inpatient Claims (MedPAR RIF), Index SNF Claims

Documentation for the Medicare claims data is provided online by the CMS contractor, Research Data Assistance Center (ResDAC) at the University of Minnesota. The following web page includes data dictionaries for the MedPAR RIF: <a href="http://www.resdac.org/cms-data/files/medpar-rif">http://www.resdac.org/cms-data/files/medpar-rif</a>

Medicare Denominator files - Documentation available at: <a href="http://www.cms.gov/Research\_Statistics-Data-and-Systems/Files-for-Order/IdentifiableDataFiles/DenominatorFile.html">http://www.cms.gov/Research\_Statistics-Data-and-Systems/Files-for-Order/IdentifiableDataFiles/DenominatorFile.html</a>

**IRF & LTCH Measure Data Sources:** The following are the specific files and links to their documentation:

Medicare Inpatient claims - standard analytical files, index PAC claims

Documentation for the Medicare claims data is provided online by ResDAC. The following web page includes data dictionaries for these files: Standard analytical files (Inpatient RIF): <a href="http://www.resdac.org/cms-data/files/ip-rif/data-documentation">http://www.resdac.org/cms-data/files/ip-rif/data-documentation</a>

Medicare Enrollment Database - Information about the Enrollment Database may be found here: <a href="http://aspe.hhs.gov/datacncl/datadir/cms.htm">http://aspe.hhs.gov/datacncl/datadir/cms.htm</a>

Medicare Denominator files - Documentation available at: <a href="http://www.cms.gov/Research-Statistics-Data-and-Systems/Files-for-Order/IdentifiableDataFiles/DenominatorFile.html">http://www.cms.gov/Research-Statistics-Data-and-Systems/Files-for-Order/IdentifiableDataFiles/DenominatorFile.html</a>

# 2.2.6 Measure Time Window

Time windows vary across settings due to variation in facility sizes across settings. The measure time window is two years in IRF and LTCH settings, and one year in the SNF setting. Specific measure time window descriptions for each setting are provided below.

*SNF Time Window:* In the SNF setting, the measure will be calculated using one year of data. All SNF stays during the one-year time window, except those that meet the exclusion criteria, will be included in the measure. For SNF residents with multiple SNF stays during the one year window, each stay is eligible for inclusion in the measure. Data from calendar year 2013 data was used to develop this PPR measure.

*Rationale:* Through the analytic work to develop this and an earlier SNF readmission measure (NQF #2510), we found one year of data to be sufficient to calculate this measure in a statistically reliable manner. This is because the reliability of a SNF's measure rate is related to the number of SNF stays included in the measure.

*IRF and LTCH Time Windows:* In the IRF and LTCH settings, the measure will be calculated using two years of data. All IRF and LTCH stays during the two-year time window, except those that meet the exclusion criteria, will be included in the measure. For patients with multiple stays during the two-year time window, each stay will be eligible for inclusion in the measure. Data from 2012-2013 were used for measure development.

Rationale: Through the analytic work to develop these and previously developed measures, we found that one year of claims data provided a somewhat limited sample size at the provider level. In order to have a more sufficient sample size, we expanded the data to include two consecutive years of claims data. In this way, the IRF and LTCH PPR measures diverge from the SNF measures which have substantially larger samples sizes compared to the IRF and LTCH settings. Pooling two years of data provides more reliable and stable estimates.

NOTE: For the purposes of public reporting, a minimum of 25 eligible stays is required.

# 2.2.7 Statistical Risk Model and Risk Adjustment Covariates

The statistical methods, including risk adjustment, were developed to harmonize with the HWR measure (NQF #1789) as well as the SNF, IRF, and LTCH all-cause readmission measures. The following section summarizes the risk adjustment approach for all PPR measures.

A hierarchical regression method using a logistic regression to predict the probability of a countable (potentially preventable, unplanned) readmission is used. The risk adjusters are predictor variables. The patient/resident characteristics related to each discharge and a marker for the specific discharging PAC provider are included in the equation. The equation is hierarchical in that both individual patient/resident characteristics are accounted for as well as the clustering of patients/residents into PAC providers. The statistical model estimates both the average predictive effect of the patient/resident characteristics across all providers and the degree to which each provider has an effect on readmissions that differs from that of the average provider. The provider effects are assumed to be randomly distributed around the average (according to a normal distribution). When computing the facility effect, hierarchical modeling accounts for the known predictors of readmissions, on average, such as patient/resident characteristics, the observed provider rate, and the number of provider stays eligible for the measure. The estimated provider effect is determined mostly by the provider's own data if the number of patient/resident discharges is relatively large (as the estimate would be relatively precise), but is adjusted toward the average if the number of patient/resident discharges is small (as that would yield an estimate of lower precision).

We used the following model:

Let  $Y_{ij}$ , denote the outcome (equal to 1 if patient/resident i is readmitted within 30 days, zero otherwise) for a patient/resident i at PAC j;  $Z_{ij}$  denotes a set of risk factors. We assume the outcome is related linearly to the covariates via a logit function with dispersion:

$$\begin{split} logit(Prob(Y_{ij} = 1)) &= \alpha_j + \beta * Z_{ij} + \epsilon_{ij} \\ \alpha_j &= \mu + \omega_j \;;\; \omega_j \sim N(0, \tau^2) \end{split} \tag{5}$$

where  $Z_{ij} = (Z_1, Z_2, ... Z_k)$  is a set of k patient-level covariates.  $\alpha_j$  represents the PAC specific intercept;  $\mu$  is the adjusted average outcome over all PAC providers; and  $\tau^2$  is the between PAC variance component and  $\varepsilon \sim N(0, \sigma^2)$  is the error term. The hierarchical logistic regression model is estimated using SAS software (PROC GLIMMIX: SAS/STAT User's Guide, SAS Institute Inc.)

NOTE: The description above refers to the method used for each measure applied to each PAC provider type and readmission window.

The estimated equation is used twice in the measure. The sum of the probabilities of readmission of all patients/residents in the measure, including both the effects of patient/resident characteristics and the provider, is the "predicted number" of readmissions after adjusting for the provider's case mix. The same equation is used without the provider effect to compute the "expected number" of potentially preventable readmissions for the same patients/residents at the average provider. The ratio of the predicted-to-expected number of readmissions is a measure of the degree to which the readmissions are higher or lower than what would otherwise be expected. This standardized risk ratio is then multiplied by the mean readmission rate for all provider stays for the measure, yielding the risk-standardized readmission rate for each provider. This estimation procedure is recalculated for each measurement period. Estimating the equations for each measurement period allows the estimated effects of the patient/resident characteristics to vary over time as medical treatment patterns change.

Risk-adjustment variables include demographic and eligibility characteristics; principal diagnoses; types of surgery or procedure from the prior short-term stay; comorbidities; length of stay and ICU/CCU utilization from the immediately prior short-term stay; and number of admissions in the year preceding the PAC admission.

The risk adjustment variables include the following:

- 1) Age/sex categories
- 2) Original reason for Medicare entitlement (age, disability or ESRD)
- 3) Surgery category if present (e.g., cardiothoracic, orthopedic), defined as in the HWR model software; the procedures are grouped using the CCS classes for ICD-9 procedures developed by AHRQ
- 4) Receiving dialysis in prior short-term stay, defined by presence of revenue code

- 5) Principal diagnosis on prior short-term claim as in the HWR measure. The ICD-9 codes are grouped clinically using the CCS for ICD-9 diagnoses developed by AHRQ.
- 6) Comorbidities from secondary diagnoses on the prior short-term claim and diagnoses from earlier short-term stays up to one year before PAC admission (these are clustered using the Hierarchical Condition Categories [HCC] groups used by CMS)

Prior Utilization Measures (vary by measure):

- 1) Length of stay in the prior short-term hospital stay (categorical to account for nonlinearity)
- 2) Prior acute ICU/CCU utilization (days) (categorical)
- 3) Count of prior short-term discharges in the prior year

PAC-Specific Risk Adjusters

- 1) IRF: Aggregates of the IRF Case-Mix Groups (CMGs) for IRF patients
- 2) LTCH: Ventilator use prolonged ventilation in LTCH (defined as ICD-9 procedure code on the index LTCH claim of 96.72, continuous invasive mechanical ventilation for 96 consecutive hours or more). We also intend to test multiple organ failure as a risk adjuster for the LTCH model.

Risk Adjustment for Sociodemographic Status (SDS):

Based on recommendations of the Consensus Standards Approval Committee, the National Quality Forum (NQF) has recently called for adjusting performance measures for sociodemographic status (SDS) when appropriate. CMS is currently conducting empirical testing under an NQF trial period to construct specific variables that capture aspects of SDS in order to account for this factor in the risk-adjustment models for the NQF-endorsed PAC readmission measures. This issue is also relevant for the potentially preventable hospital readmission measures that are currently under development. In addition, work being conducted by the Assistant Secretary for Planning and Evaluation on SDS risk adjustment per the IMPACT Act may provide additional direction on this issue.

## 2.2.8 Measure Calculation Algorithm

We are developing 4 PPR measures; each is specific to a single PAC provider type and readmission window. Because the overall calculation algorithms and logic are aligned for the set of PPR PAC measures, we describe these technical details for all measures below, rather than duplicating similar information. It is important to clarify that each measure is specific to a single PAC provider type; we do not pool PAC patients/residents across settings in the measure calculation.

The Medicare PAC claims are matched to prior acute hospital stays, hospital stays post-PAC discharge, and patient/resident eligibility data to determine which stays remain in the measure (i.e. not excluded per the exclusions described above) and which have potentially preventable, unplanned readmissions.

The measures are calculated according to the following steps:

- Step 1: Identify patients/residents meeting the denominator (measure inclusion) criteria.
- Step 2: Identify patients/residents meeting the numerator (unplanned PPR) criteria taking into account the planned readmission algorithm.
- Step 3: Identify presence or absence of risk adjustment variables for each patient/resident.
- Step 4: Calculate the predicted and expected number of readmissions for each PAC provider using hierarchical logistic regression model.

The predicted number of readmissions for each PAC provider for each measure is calculated as the sum of the predicted probability of readmission for each patient/resident included in the measure discharged from the provider, including the provider-specific effect. The model specific risk standardized readmission ratio for each PAC provider associated with each PPR measure is calculated as follows.

To calculate the predicted number of readmissions  $pred_j$  for index PAC provider stays at provider, we used

$$pred_{j} = \sum logit^{-1}(\mu + \omega_{i} + \beta*Z_{ij})$$
 (6)

where the sum is over all stays in provider<sub>j</sub>, and  $\omega_i$  is the random intercept. To calculate the expected number  $exp_j$  use

$$exp_{j} = \Sigma logit^{-1} (\mu + \beta * Z_{ij})$$
 (7)

Then, as a measure of excess or reduced readmissions among index stays at PAC provider<sub>j</sub>, calculate the provider-wide standardized risk ratio, SRR<sub>j</sub>, as

$$SRR_{i} = pred_{i}/exp_{i}$$
 (8)

Step 5: Calculate the risk-standardized PAC potentially preventable readmission rate.

The value obtained from equation (4) above, the  $SRR_j$ , is the PAC provider-wide standardized risk ratio for provider<sub>j</sub>. To aid interpretation, the provider-wide standardized risk ratio,  $SRR_j$ , is then multiplied by the overall national raw readmission rate for all provider stays,  $\bar{Y}$ , to produce the provider-wide risk-standardized readmission rate ( $RSRR_j$ ).

$$RSRR_{j} = SRR_{j} * \bar{Y}$$
 (9)

# 2.2.9 Measure Results

We present measure results for the set of PPR measures in the **Appendix 2, Table 2-8** and **Figures 2-2 and 2-3**. These appendices include the full risk adjustment model results along with distributions of the unadjusted and risk-standardized PPR rates for facilities.

# SECTION 3 MEASURES AFFECTING THE FY 2020 PAYMENT DETERMINATION AND SUBSEQUENT YEARS

# 3.1 Drug Regimen Review Conducted with Follow-Up for Identified Issues- Post Acute Care (PAC) Skilled Nursing Facility (SNF) Quality Reporting Program (QRP)

# 3.1.1 Measure Description

Under sections 1899B(a)(2)(E)(i)(III) and 1899B(c)(1)(C) of the Act requires the Secretary to specify a quality measure to address the medication reconciliation domain for IRFs, LTCHs and SNFs by October 1, 2018 and for HHAs by January 1, 2017. We are proposing to adopt the quality measure, Drug Regimen Review Conducted with Follow-Up for Identified Issues- Post Acute Care (PAC) Skilled Nursing Facility (SNF) Quality Reporting Program (QRP), for the SNF QRP as a resident assessment-based, cross-setting quality measure to meet the SNF requirements with data collection beginning October 1, 2018 for FY 2020 payment determinations and subsequent years.

This proposed measure assesses whether PAC providers were responsive to potential or actual clinically significant medication issue(s) when such issues were identified. Specifically, the proposed quality measure reports the percentage of patient/resident stays in which a drug regimen review was conducted at the time of admission and timely follow-up with a physician occurred each time potential clinically significant medication issues were identified throughout that stay.

Additionally, for this proposed quality measure, drug regimen review is defined as the review of all medications or drugs the patient/resident is taking to identify any potentially clinically significant medication issues. This proposed quality measure utilizes both the processes of medication reconciliation and a drug regimen review, in the event an actual or potential medication issue occurred. The proposed measure informs whether the PAC facility identified and addressed each clinically significant medication issue and if the facility responded or addressed the medication issue in a timely manner. Of note, drug regimen review in PAC settings is generally considered to include medication reconciliation and review of the patient's drug regimen to identify potential clinically significant medication issues. This measure is applied uniformly across the PAC settings.

For SNFs, this measure applies to resident stays covered by Medicare Part A. In IRFs, this measure is limited to Medicare (Part A and Part C) patients. In LTCHs, this measure includes all patients.

## 3.1.2 Purpose/Rationale for the Quality Measure

This proposed measure assesses whether PAC providers were responsive to potential or actual clinically significant medication issue(s) when such issues were identified. Specifically, the proposed quality measure reports the percentage of patient/resident stays in which a drug

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<sup>&</sup>lt;sup>65</sup>Institute of Medicine. Preventing Medication Errors. Washington DC: National Academies Press; 2006.
<sup>66</sup>Ibid.

regimen review was conducted at the time of admission and timely follow-up with a physician occurred each time potential clinically significant medication issues were identified throughout that stay.

The performance of timely medication reconciliation is valuable to the process of drug regimen review. Preventing and responding to ADEs is of critical importance as ADEs account for significant increases in health services utilization and costs, <sup>67, 68, 69</sup> including subsequent emergency room visits and re-hospitalizations. Annual health care costs in the United States are estimated at \$3.5 billion, resulting in 7,000 deaths annually. <sup>71,72</sup>

Medication reconciliation is a process of reviewing an individual's complete and current medication list and drug regimen review is included in that process. Medication reconciliation is a recognized process for reducing the occurrence of medication discrepancies that may lead to Adverse Drug Events (ADEs). 73 Medication discrepancies occur when there is conflicting information documented in the medical records. The World Health Organization regards medication reconciliation as a standard operating protocol necessary to reduce the potential for ADEs that cause harm to patients. Medication reconciliation is an important patient safety process that addresses medication accuracy during transitions in patient care and in identifying preventable ADEs. 74 The Joint Commission added medication reconciliation to its list of National Patient Safety Goals (2005), suggesting that medication reconciliation is an integral component of medication safety. 75 The Society of Hospital Medicine published a statement in agreement of the Joint Commission's emphasis and value of medication reconciliation as a patient safety goal. 76 There is universal agreement that medication reconciliation directly

<sup>67</sup> Institute of Medicine. Preventing Medication Errors. Washington DC: National Academies Press; 2006.

<sup>68</sup> Jha AK, Kuperman GJ, Rittenberg E, et al. Identifying hospital admissions due to adverse drug events using a computer-based monitor. Pharmacoepidemiol Drug Saf. 2001;10(2):113-119.

<sup>69</sup> Hohl CM, Nosyk B, Kuramoto L, et al. Outcomes of emergency department patients presenting with adverse drug events. Ann Emerg Med. 2011;58:270–279.

<sup>70</sup> Kohn LT, Corrigan JM, Donaldson MS. To Err Is Human: Building a Safer Health System Washington, DC: National Academies Press; 1999.

<sup>71</sup> Greenwald, J. L., Halasyamani, L., Greene, J., LaCivita, C., et al. (2010). Making inpatient medication reconciliation patient centered, clinically relevant and implementable: a consensus statement on key principles and necessary first steps. Journal of Hospital Medicine, 5(8), 477-485.

<sup>72</sup> Phillips, David P.; Christenfeld, Nicholas; and Glynn, Laura M. Increase in US Medication-Error Deaths between 1983 and 1993. The Lancet. 351:643–644, 1998.

<sup>73</sup> Institute of Medicine. Preventing Medication Errors. Washington DC: National Academies Press; 2006.

<sup>74</sup> Leotsakos A., et al. Standardization in patient safety: the WHO High 5s project. Int J Qual Health Care. 2014;26(2):109-116.

<sup>75</sup> The Joint Commission. 2016 Long Term Care: National Patient Safety Goals Medicare/Medicaid Certification-based Option. (NPSG.03.06.01).

<sup>76</sup> Greenwald, J. L., Halasyamani, L., Greene, J., LaCivita, C., et al. (2010). Making inpatient medication reconciliation patient centered, clinically relevant and implementable: a consensus statement on key principles and necessary first steps. Journal of Hospital Medicine, 5(8), 477-485.

addresses patient safety issues that can result from medication miscommunication and unavailable or incorrect information. 77,78,79

Medication errors include the duplication of medications, delivery of an incorrect drug, inappropriate drug omissions, or errors in the dosage, route, frequency, and duration of medications. Medication errors are one of the most common types of medical error and can occur at any point in the process of ordering and delivering a medication. Medication errors have the potential to result in an ADE. 80,81,82,83,84,85 Inappropriately prescribed medications are also considered a major healthcare concern in the United States for the elderly population, with costs of roughly \$7.2 billion annually. 86

There is strong evidence that medication discrepancies occur during transfers from acute care facilities to post-acute care facilities. Discrepancies occur when there is conflicting information documented in the medical records. Almost one-third of medication discrepancies have the potential to cause patient harm. <sup>87</sup> An estimated fifty percent of patients experienced a clinically important medication error after hospital discharge in an analysis of two tertiary care academic hospitals. <sup>88</sup>

Medication reconciliation has been identified as an area for improvement during transfer from the acute care facility to the receiving post-acute care facility. Post-acute care facilities report gaps in medication information between the acute care hospital and the receiving post-

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<sup>&</sup>lt;sup>77</sup> Leotsakos A., et al. Standardization in patient safety: the WHO High 5s project. Int J Qual Health Care. 2014:26(2):109-116.

<sup>78</sup> The Joint Commission. 2016 Long Term Care: National Patient Safety Goals Medicare/Medicaid Certification-based Option. (NPSG.03.06.01).

<sup>79</sup> IHI. Medication Reconciliation to Prevent Adverse Drug Events [Internet]. Cambridge, MA: Institute for Healthcare Improvement; [cited 2016 Jan 11]. Available from: <a href="http://www.ihi.org/topics/adesmedicationreconciliation/Pages/default.aspx">http://www.ihi.org/topics/adesmedicationreconciliation/Pages/default.aspx</a>.

<sup>80</sup> Institute of Medicine. To err is human: building a safer health system. Washington, DC: National Academies Press; 2000.

<sup>81</sup> Lesar TS, Briceland L, Stein DS. Factors related to errors in medication prescribing. JAMA. 1997:277(4): 312-317.

<sup>82</sup> Bond CA, Raehl CL, & Franke T. Clinical pharmacy services, hospital pharmacy staffing, and medication errors in United States hospitals. Pharmacotherapy. 2002:22(2): 134-147.

<sup>83</sup> Bates DW, Cullen DJ, Laird N, Petersen LA, Small SD, et al. Incidence of adverse drug events and potential adverse drug events. Implications for prevention. JAMA. 1995:274(1): 29-34.

<sup>84</sup> Barker KN, Flynn EA, Pepper GA, Bates DW, & Mikeal RL. Medication errors observed in 36 health care facilities. JAMA. 2002: 162(16):1897-1903.

<sup>85</sup> Bates DW, Boyle DL, Vander Vliet MB, Schneider J, & Leape L. Relationship between medication errors and adverse drug events. J Gen Intern Med. 1995:10(4): 199-205.

<sup>&</sup>lt;sup>86</sup> Fu, Alex Z., et al. "Potentially inappropriate medication use and healthcare expenditures in the US community-dwelling elderly." Medical care 45.5 (2007): 472-476.

<sup>87</sup> Wong, Jacqueline D., et al. "Medication reconciliation at hospital discharge: evaluating discrepancies." Annals of Pharmacotherapy 42.10 (2008): 1373-1379.

<sup>88</sup> Kripalani S, Roumie CL, Dalal AK, et al. Effect of a pharmacist intervention on clinically important medication errors after hospital discharge: A randomized controlled trial. Ann Intern Med. 2012:157(1):1-10.

acute care setting when performing medication reconciliation. <sup>89,90</sup> Hospital discharge has been identified as a particularly high risk point in time, with evidence that medication reconciliation identifies high levels of discrepancy. <sup>91,92,93,94,95,96</sup> Also, there is evidence that medication reconciliation discrepancies occur throughout the patient stay. <sup>97</sup>, <sup>98</sup> For older patients, who may have multiple comorbid conditions and thus multiple medications, transitions between acute and post-acute care settings can be further complicated, <sup>99</sup> and medication reconciliation and patient knowledge (medication literacy) can be inadequate post-discharge. <sup>100</sup> The proposed quality measure, Drug Regimen Review Conducted with Follow-Up for Identified Issues-PAC SNF QRP, provides an important component of care coordination for PAC settings and would affect a large proportion of the Medicare population who transfer from hospitals into PAC services each year. For example, in 2013, 1.7 million Medicare FFS beneficiaries had SNF stays, 338,000 beneficiaries had IRF stays, and 122,000 beneficiaries had LTCH stays. <sup>101</sup>

#### 3.1.3 Denominator

The denominator is the number of stays during the SNF, IRF, or LTCH reporting period. Specific denominator definitions for each setting are provided below.

**SNF Denominator:** The denominator is the number of stays in the selected time window for SNF residents with a SNF PPS Part A Discharge Assessment (A0310H = 1) during the reporting period. A stay is defined as the time period from resident admission or reentry to the

89 Gandara, Esteban, et al. "Communication and information deficits in patients discharged to rehabilitation facilities: an evaluation of five acute care hospitals." Journal of Hospital Medicine 4.8 (2009): E28-E33.

<sup>90</sup> Gandara, Esteban, et al. "Deficits in discharge documentation in patients transferred to rehabilitation facilities on anticoagulation: results of a system wide evaluation." Joint Commission Journal on Quality and Patient Safety 34.8 (2008): 460-463.

<sup>91</sup> Coleman EA, Smith JD, Raha D, Min SJ. Post hospital medication discrepancies: prevalence and contributing factors. Arch Intern Med. 2005 165(16):1842–1847.

<sup>92</sup> Wong JD, Bajcar JM, Wong GG, et al. Medication reconciliation at hospital discharge: evaluating discrepancies. Ann Pharmacother. 2008 42(10):1373–1379.

<sup>93</sup> Hawes EM, Maxwell WD, White SF, Mangun J, Lin FC. Impact of an outpatient pharmacist intervention on medication discrepancies and health care resource utilization in post hospitalization care transitions. Journal of Primary Care & Community Health. 2014; 5(1):14-18.

<sup>94</sup> Foust JB, Naylor MD, Bixby MB, Ratcliffe SJ. Medication problems occurring at hospital discharge among older adults with heart failure. Research in Gerontological Nursing. 2012, 5(1): 25-33.

<sup>95</sup> Pherson EC, Shermock KM, Efird LE, et al. Development and implementation of a post discharge home-based medication management service. Am J Health Syst Pharm. 2014; 71(18): 1576-1583.

<sup>96</sup> Pronovosta P, Weasta B, Scwarza M, et al. Medication reconciliation: a practical tool to reduce the risk of medication errors. J Crit Care. 2003; 18(4): 201-205.

<sup>97</sup> Bates DW, Cullen DJ, Laird N, Petersen LA, Small SD, et al. Incidence of adverse drug events and potential adverse drug events. Implications for prevention. JAMA. 1995:274(1): 29-34.

<sup>&</sup>lt;sup>98</sup> Himmel, W., M. Tabache, and M. M. Kochen. "What happens to long-term medication when general practice patients are referred to hospital? "European journal of clinical pharmacology 50.4 (1996): 253-257.

<sup>&</sup>lt;sup>99</sup> Chhabra, P. T., et al. (2012). "Medication reconciliation during the transition to and from long-term care settings: a systematic review." Res Social Adm Pharm 8(1): 60-75.

<sup>100</sup> Kripalani S, Roumie CL, Dalal AK, et al. Effect of a pharmacist intervention on clinically important medication errors after hospital discharge: A randomized controlled trial. Ann Intern Med. 2012:157(1):1-10.

<sup>101</sup> March 2015 Report to the Congress: Medicare Payment Policy. Medicare Payment Advisory Commission; 2015.

facility (identified by a 5-day PPS assessment) to discharge, which may be an OBRA discharge or a SNF PPS Part A Discharge.

**LTCH Denominator**: The denominator is the number of patient stays with a discharge or expired assessment (A0250=10, 11, 12) during the reporting period.

*IRF Denominator:* The denominator is the number of Medicare patient stays\* (Part A or Part C) during the reporting period.

\*IRF-PAI data are submitted only for Medicare patients (Part A and Part C).

#### Denominator Exclusions

This measure has no denominator exclusions for IRF, LTCH, and SNF.

#### 3.1.4 Numerator

Number of stays in the denominator where the medical record contains documentation of a drug regimen review conducted at admission with all potential clinically significant medication issues identified during the course of care and followed-up with a physician or physician designee.

Specific numerator definitions for each setting are provided below.

*SNF Numerator*: The numerator is the number of short-stay residents with an MDS 3.0 assessment during the selected time window for which all of the following are each true:

- 1) The facility conducted a drug regimen review at the admission (N2001= [0,1]) or resident is not taking any medications (N2001= [9]); and
- 2) If potential clinically significant medication issues were identified at the admission (N2001 = [1]), then the facility contacted a physician (or physician-designee) by midnight of the next calendar day and completed prescribed/recommended actions in response to the identified issues (N2003= [1]); and
- 3) The facility contacted a physician (or physician-designee) and completed prescribed/recommended actions by midnight of the next calendar day each time potential clinically significant medication issues were identified since the admission (N2005 = [1]) or no potential clinically significant medications issues were identified since the admission (N2005 = [9]). This condition is evaluated at discharge.

**LTCH Numerator:** The numerator is the number of stays for which the LTCH CARE Data Set indicated all of the following are each true:

- 1) The facility conducted a drug regimen review at the admission (N2001=[0,1]) or patient is not taking any medications (N2001=[9]); and
- 2) If potential clinically significant medication issues were identified at the admission (N2001 = [1]), then the facility contacted a physician (or physician-designee) by

- midnight of the next calendar day and completed prescribed/recommended actions in response to the identified issues (N2003=[1]); and
- 3) The facility contacted a physician (or physician-designee) and completed prescribed/recommended actions by midnight of the next calendar day each time potential clinically significant medication issues were identified since the admission (N2005 = [1]) or no potential clinically significant medications issues were identified since the admission (N2005 = [9]).

*IRF Numerator:* The numerator is the number of stays for which the IRF PAI indicated all of the following are each true:

- 1) The facility conducted a drug regimen review at the admission (N2001=[0,1]) or patient is not taking any medications (N2001=[9]); and
- 2) If potential clinically significant medication issues were identified at the admission (N2001 = [1]), then the facility contacted a physician (or physician-designee) by midnight of the next calendar day and completed prescribed/recommended actions in response to the identified issues (N2003= [1]); and
- 3) The facility contacted a physician (or physician-designee) and completed prescribed/recommended actions by midnight of the next calendar day each time potential clinically significant medication issues were identified since the admission (N2005 = [1]) or no potential clinically significant medications issues were identified since the admission (N2005 = [9]).

Please note that if data is missing on any of the three items used to calculate the numerator of the measure (specifically, (N2001=[-] or N2003=[-] or N2005=[-])), the patient/resident will not be included in the numerator count though they will continue to be counted in the denominator, assuming all denominator criteria for that patient/resident have been met.

## 3.1.5 Items Included in the Quality Measure

See **Appendix 3, Table 3-1** for a summary of the setting specific language used to describe the resident or patient within the PAC setting. There are no other differences in the content language within each Drug Regimen Review quality measure item.

## N2001. Drug Regimen Review Item

Did a complete drug regimen review identify potential clinically significant medication issues?

- **0.** No No issues found during review
- **1. Yes** Issues found during review
- **9.** NA Patient/Resident is not taking any medications

# **N2003 Medication Follow-up Item**

Did the facility contact a physician (or physician-designee) by midnight of the next calendar day and complete prescribed/recommended actions in response to the identified potential clinically significant medication issues?

- 0. No
- 1. Yes

#### N2005. Medication Intervention Item

Did the facility contact and complete physician (or physician-designee) prescribed/recommended actions by midnight of the next calendar day each time potential clinically significant medication issues were identified since the Admission?

- 0. No
- 1. Yes
- **9.** NA There were no potential clinically significant medication issues identified since Admission or patient/resident is not taking any medications.

#### 3.1.6 Risk Adjustment

This measure is not risk-adjusted or stratified.

#### 3.1.7 Quality Measure Calculation Algorithm

The following steps are used to calculate the measure:

Step 1: Calculate the denominator count (see Section 3.1.3 for details):

In the SNF setting, identify SNF residents with a PPS Part A Discharge (A0310H=1). Count the number of SNF stays (resident admission or reentry to the facility to discharge, which may be an OBRA discharge or a SNF PPS Part A Discharge) among these residents.

In the LTCH setting, calculate the number of patient stays with a discharge or expired assessment (A0250=10, 11, 12),

In the IRF setting, calculate the number of Medicare (Part A or Part C) patient stays.

Step 2: Calculate the numerator count (see Section 3.1.4 for details):

In the SNF setting, calculate the total number short-stay resident stays in the denominator where the medical record contains documentation of a drug regimen review conducted at: 1) admission and 2) throughout the stay, with all potential clinically significant medication issues identified during the course of care and followed up with a physician or physician designee by midnight of the next calendar day.

In the LTCH setting, calculate the total number of patient stays whose LTCH-CARE Data Set assessment indicates that the medical record contains documentation of a drug regimen review conducted at:

1) Admission and 2) throughout the stay, with all potential clinically significant medication issues identified during the course of care and followed up with a physician or physician designee by midnight of the next calendar day.

In the IRF setting, calculate the total number of patient stays whose IRF-PAI assessment indicates that the medical record contains documentation of a drug regimen review conducted at: 1) Admission and 2) throughout the stay, with all potential clinically significant medication issues identified during the course of care and followed up with a physician or physician designee by midnight of the next calendar day.

# Step 3: Calculate the facility's observed score:

Divide the facility's numerator count by its denominator count to obtain the facility's observed score; that is, divide the result of step 2 by the result of step 1. [This page intentionally left blank.]

# APPENDIX 1 DISCHARGE TO COMMUNITY- POST ACUTE CARE (PAC) SKILLED NURSING FACILITY (SNF) QUALITY REPORTING PROGRAM (QRP)

- Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled Nursing Facility Quality Reporting Program, 2013
- Table 1-2. Skilled Nursing Facility: Facility-Level Observed and Risk-Standardized Discharge to Community Rates, 2013
- Figure 1-1. Skilled Nursing Facility: Facility-Level Observed and Risk-Standardized Discharge to Community Rates, 2013

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled
Nursing Facility Quality Reporting Program, 2013

Number of beneficiaries included in the model = 1,846,070

Observed number (percent) of beneficiaries in the sample who were discharged to community = 817,562 (44.3%).

Model c-statistic = 0.744

Based on Medicare fee-for-service claims data from CY 2013. These model estimates only apply to CY 2013 SNF data. We will re-estimate the regression models for each measurement period to allow the estimated effects of resident characteristics to vary over time.

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
Intercept			2.136	0.013	<.0001			
Age and Sex Groupings (Reference: Female, age 65-69 years)								
Male, age 18-35 years	1,605	0.09	-0.204	0.059	0.0005	0.816	0.727	0.915
Male, age 35-44 years	4,960	0.27	-0.082	0.034	0.0151	0.922	0.863	0.984
Male, age 45-54 years	20,940	1.13	-0.171	0.018	<.0001	0.842	0.813	0.873
Male, age 55-59 years	22,834	1.24	-0.216	0.017	<.0001	0.806	0.779	0.834
Male, age 60-64 years	30,322	1.64	-0.275	0.015	<.0001	0.759	0.737	0.783
Male, age 65-69 years	81,023	4.39	-0.064	0.011	<.0001	0.938	0.918	0.958
Male, age 70-74 years	85,445	4.63	-0.158	0.011	<.0001	0.854	0.836	0.872
Male, age 75-79 years	104,634	5.67	-0.217	0.011	<.0001	0.805	0.789	0.822
Male, age 80-84 years	124,518	6.75	-0.301	0.010	<.0001	0.740	0.725	0.755
Male, age 85-89 years	124,014	6.72	-0.429	0.010	<.0001	0.651	0.638	0.664
Male, age 90-94 years	72,100	3.91	-0.607	0.012	<.0001	0.545	0.533	0.558
Male, age ≥ 95 years	20,316	1.10	-0.877	0.018	<.0001	0.416	0.402	0.431
Female, age 18-35 years	1,386	0.08	-0.134	0.063	0.0344	0.875	0.773	0.990
Female, age 35-44 years	4,312	0.23	-0.154	0.036	<.0001	0.857	0.799	0.920
Female, age 45-54 years	18,377	1.00	-0.118	0.019	<.0001	0.889	0.856	0.922
Female, age 55-59 years	22,173	1.20	-0.118	0.017	<.0001	0.889	0.859	0.920
Female, age 60-64 years	33,031	1.79	-0.152	0.015	<.0001	0.859	0.834	0.884

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled
Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
Female, age 70-74 years	125,285	6.79	-0.040	0.010	<.0001	0.961	0.942	0.980
Female, age 75-79 years	164,744	8.92	-0.114	0.010	<.0001	0.893	0.876	0.910
Female, age 80-84 years	211,135	11.44	-0.238	0.009	<.0001	0.788	0.774	0.803
Female, age 85-89 years	239,344	12.97	-0.398	0.009	<.0001	0.672	0.660	0.684
Female, age 90-94 years	165,658	8.97	-0.612	0.010	<.0001	0.542	0.532	0.553
Female, age ≥ 95 years	63,300	3.43	-0.908	0.012	<.0001	0.404	0.394	0.413
Original Reason for Entitlement						_		
Age $\geq$ 65 at SNF admission and original reason for entitlement was disability	243,141	13.17	-0.224	0.005	<.0001	0.799	0.791	0.808
Age $\geq$ 65 at SNF admission and original reason for entitlement was ESRD, or ESRD and disability	8,291	0.45	-0.206	0.029	<.0001	0.814	0.769	0.862
Principal Diagnosis Clinical Classifications Software (CCS) Groupings ba Procedures (10, 255, 256, 258) and Fetal and Neonate Conditions (218-224		Acute Stay	(Reference: O	steo and	Infective	Arthritis (20	1, 203, 204, 20	6),
Infections (TB, Bacterial, HIV, Other) (1, 3, 4, 5, 7, 8, 9)	6,234	0.34	-1.033	0.030	<.0001	0.356	0.335	0.377
Septicemia (except in labor) (2)	148,235	8.03	-1.081	0.015	<.0001	0.339	0.330	0.350
Cancer: Cancer of Head and Neck (11)	771	0.04	-1.209	0.084	<.0001	0.299	0.253	0.352
Cancer: Stomach, Colon (12, 13, 14)	6,794	0.37	-0.665	0.030	<.0001	0.514	0.485	0.545
Cancer: Cancer of Rectum and Anus (15)	1,607	0.09	-1.146	0.055	<.0001	0.318	0.286	0.354
Cancer: Other GI (16, 17, 18)	1,488	0.08	-0.911	0.056	<.0001	0.402	0.360	0.449
Cancer: Cancer of Bronchus; Lung; Cancer; Other Respiratory and Intrathoracic (19, 20)	2,205	0.12	-1.239	0.050	<.0001	0.290	0.263	0.319
Cancer: Bone and Melanoma (21, 22, 23, 26, 28, 29, 30, 31, 36)	1,597	0.09	-1.015	0.056	<.0001	0.362	0.325	0.404
Cancer: Breast and Female (24, 25, 27)	1,894	0.10	-1.035	0.056	<.0001	0.355	0.318	0.396
Cancer: Bladder and Kidney (32, 33, 34)	3,003	0.16	-1.054	0.045	<.0001	0.349	0.319	0.381
Cancer: Cancer of Brain and Nervous System (35)	677	0.04	-1.652	0.087	<.0001	0.192	0.162	0.227

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled
Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
Cancer: Leukemia and Lymphoma (37, 38, 39, 40, 41, 43, 45)	1,081	0.06	-1.870	0.070	<.0001	0.154	0.134	0.177
Cancer: Secondary Malignancies (42)	2,689	0.15	-1.383	0.044	<.0001	0.251	0.230	0.273
Cancer: Misc. Neoplasm (44, 46, 47, 167)	3,806	0.21	-0.925	0.037	<.0001	0.397	0.369	0.426
Thyroid Disorders (48)	1,061	0.06	-1.050	0.067	<.0001	0.350	0.307	0.399
Diabetes Mellitus with or without Complication (49, 50)	26,434	1.43	-0.963	0.017	<.0001	0.382	0.369	0.395
Other Endocrine Disorders; Disorders of Lipid Metabolism (51, 53)	5,120	0.28	-0.860	0.031	<.0001	0.423	0.398	0.450
Nutritional Deficiencies (52)	1,241	0.07	-1.390	0.065	<.0001	0.249	0.219	0.283
Gout and Other Crystal Arthropathies (54)	1,477	0.08	-0.585	0.056	<.0001	0.557	0.499	0.622
Fluid/Electrolyte Disorders (55)	30,804	1.67	-1.141	0.016	<.0001	0.320	0.309	0.330
COPD, Asthma, and Cystic Fibrosis (56, 127, 128)	47,666	2.58	-1.252	0.015	<.0001	0.286	0.278	0.294
Blood Disorders (Immune, Sickle Cell) (57, 61, 62, 63, 64)	2,942	0.16	-1.175	0.042	<.0001	0.309	0.284	0.335
Other Nutritional, Endocrine, and Metabolic Disorders (58)	5,266	0.29	-1.250	0.032	<.0001	0.286	0.269	0.305
Iron Deficiency and Other Anemia (59)	13,149	0.71	-1.431	0.023	<.0001	0.239	0.229	0.250
Acute Posthemorrhagic Anemia (60)	3,846	0.21	-1.072	0.037	<.0001	0.342	0.318	0.368
Meningitis and Encephalitis (76, 77, 78)	1,962	0.11	-0.798	0.049	<.0001	0.450	0.409	0.496
Parkinson's Disease (79)	2,013	0.11	-0.939	0.048	<.0001	0.391	0.356	0.429
Multiple Sclerosis (80)	974	0.05	-0.758	0.070	<.0001	0.469	0.409	0.537
Other Hereditary and Degenerative Nervous System Conditions; Paralysis (81, 82)	4,714	0.26	-0.986	0.033	<.0001	0.373	0.350	0.398
Epilepsy, Convulsions (83)	11,465	0.62	-1.044	0.023	<.0001	0.352	0.336	0.368
Migraine and Eye Conditions (84, 86, 87, 88, 89, 90, 91, 92, 93, 94)	4,056	0.22	-0.542	0.035	<.0001	0.582	0.543	0.623
Coma, Stupor, and Brain Damage (85)	1,667	0.09	-1.100	0.055	<.0001	0.333	0.299	0.370

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled
Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
Other Nervous System Disorders (95)	23,059	1.25	-0.912	0.018	<.0001	0.402	0.388	0.416
Heart Valve Disorders (96)	12,823	0.69	-0.998	0.026	<.0001	0.369	0.350	0.388
Cardiomyopathy; Coronary Atherosclerosis and other Heart Disease (97, 101)	15,443	0.84	-0.963	0.022	<.0001	0.382	0.366	0.399
Essential Hypertension; Hypertension with Complications and Secondary Hypertension (98, 99)	15,376	0.83	-1.016	0.021	<.0001	0.362	0.348	0.377
Acute MI; Cardiac Arrest and Ventricular Fibrillation (100, 107)	32,701	1.77	-1.141	0.016	<.0001	0.319	0.309	0.330
Nonspecific Chest Pain; Other and Ill-defined Heart Disease (102, 104)	5,685	0.31	-1.117	0.031	<.0001	0.327	0.308	0.348
Pulmonary Heart Disease (103)	11,413	0.62	-0.900	0.023	<.0001	0.406	0.389	0.425
Conduction Disorders; Cardiac Dysrhythmias (105, 106)	42,185	2.29	-1.005	0.015	<.0001	0.366	0.356	0.377
Congestive Heart Failure (108)	87,262	4.73	-1.222	0.013	<.0001	0.295	0.287	0.303
Acute Cerebrovascular Disease (109)	66,476	3.60	-1.082	0.014	<.0001	0.339	0.330	0.348
Occlusion or Stenosis of Precerebral Arteries (110)	2,238	0.12	-0.890	0.048	<.0001	0.411	0.374	0.451
Other and Ill-defined Cerebrovascular Disease (111)	1,165	0.06	-1.056	0.067	<.0001	0.348	0.305	0.397
Transient Cerebral Ischemia (112)	8,406	0.46	-0.731	0.025	<.0001	0.481	0.458	0.506
Late Effects of Cerebrovascular Disease (113)	2,192	0.12	-0.924	0.048	<.0001	0.397	0.361	0.436
Peripheral and Visceral Atherosclerosis (114)	10,313	0.56	-1.055	0.025	<.0001	0.348	0.332	0.365
Aortic, Peripheral, and Visceral Artery Aneurysms (115)	3,829	0.21	-0.848	0.038	<.0001	0.428	0.397	0.461
Aortic and Peripheral Arterial Embolism or Thrombosis (116)	2,521	0.14	-1.182	0.045	<.0001	0.307	0.281	0.335
Other Circulatory Disease (117)	9,735	0.53	-0.792	0.024	<.0001	0.453	0.432	0.475
Phlebitis, Thrombophlebitis and Thromboembolism (118)	10,544	0.57	-1.044	0.023	<.0001	0.352	0.336	0.369
Vein Disease and Lymphadenitis (119, 120, 121, 247)	3,504	0.19	-1.034	0.038	<.0001	0.356	0.330	0.383
Pneumonia (except that caused by Tuberculosis or Sexually Transmitted Disease) (122)	88,591	4.80	-1.234	0.013	<.0001	0.291	0.284	0.299

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled
Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
Influenza (123)	5,717	0.31	-0.829	0.030	<.0001	0.436	0.412	0.463
Tonsillitis and Teeth and Jaw Disorders (124, 136, 137)	1,137	0.06	-1.431	0.069	<.0001	0.239	0.209	0.274
Acute Bronchitis; Other Upper Respiratory Infections (125, 126)	4,304	0.23	-0.914	0.034	<.0001	0.401	0.375	0.428
Aspiration Pneumonitis, Food/Vomitus (129)	29,126	1.58	-1.556	0.019	<.0001	0.211	0.203	0.219
Pleurisy, Pneumothorax, Pulmonary Collapse (130)	6,465	0.35	-1.140	0.030	<.0001	0.320	0.302	0.339
Respiratory Failure, Insufficiency, Arrest (Adult) (131)	31,666	1.72	-1.299	0.017	<.0001	0.273	0.264	0.282
Other Lung Disease (132, 133, 134)	5,762	0.31	-1.241	0.031	<.0001	0.289	0.272	0.307
Intestinal Infection (135)	17,383	0.94	-1.115	0.020	<.0001	0.328	0.316	0.341
Esophageal Disorders (138)	4,360	0.24	-1.179	0.035	<.0001	0.308	0.287	0.330
Gastroduodenal Ulcer (except Hemorrhage) (139)	1,885	0.10	-1.007	0.050	<.0001	0.365	0.331	0.403
Gastritis and Duodenitis; Other Disorders of Stomach and Duodenum (140, 141)	6,562	0.36	-1.131	0.029	<.0001	0.323	0.305	0.342
Appendicitis and Other Appendiceal Conditions (142)	1,194	0.06	-0.738	0.063	<.0001	0.478	0.422	0.541
Abdominal Hernia (143)	7,669	0.42	-0.867	0.028	<.0001	0.420	0.398	0.444
Ulcerative Colitis and Diverticulitis (144, 146, 148)	13,941	0.76	-0.988	0.021	<.0001	0.372	0.357	0.388
Intestinal Obstruction without Hernia (145)	17,922	0.97	-1.022	0.020	<.0001	0.360	0.346	0.374
Anal and Rectal Conditions (147)	2,177	0.12	-1.078	0.048	<.0001	0.340	0.310	0.374
Biliary Tract Disease (149)	11,813	0.64	-0.957	0.023	<.0001	0.384	0.367	0.402
Other Liver Disease (6, 150, 151)	8,180	0.44	-1.574	0.030	<.0001	0.207	0.195	0.220
Pancreatic Disorders (not Diabetes) (152)	4,800	0.26	-0.872	0.032	<.0001	0.418	0.392	0.446
Gastrointestinal Hemorrhage (153)	26,744	1.45	-1.234	0.017	<.0001	0.291	0.281	0.301
Noninfectious Gastroenteritis; Other Gastrointestinal Disorders (154, 155)	13,821	0.75	-1.031	0.021	<.0001	0.357	0.342	0.372

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
Nephritis; Nephrosis; Renal Sclerosis; Acute and Unspecified Renal Failure (156, 157)	61,399	3.33	-1.189	0.014	<.0001	0.305	0.296	0.313
Chronic Renal Failure (158)	803	0.04	-1.250	0.085	<.0001	0.287	0.243	0.338
Urinary Tract Infections (159)	73,182	3.96	-1.236	0.014	<.0001	0.291	0.283	0.298
Calculus of Urinary Tract (160)	1,834	0.10	-1.096	0.053	<.0001	0.334	0.302	0.371
Other Diseases of Kidney and Ureters; Other Diseases of Bladder and Urethra(161, 162)	2,547	0.14	-1.110	0.045	<.0001	0.329	0.302	0.360
Genitourinary Symptoms and Ill-defined Conditions (163)	2,272	0.12	-1.218	0.048	<.0001	0.296	0.269	0.325
Prostate and Other Male Genital (164, 165, 166)	2,700	0.15	-0.981	0.043	<.0001	0.375	0.345	0.408
Female Reproductive (168, 169, 170, 171, 172, 173, 174, 175, 176, 177, 178, 179, 180, 181, 182, 183, 184, 185, 186, 187, 188, 189, 190, 191, 192, 193, 194, 195, 196)	1,406	0.08	-1.100	0.061	<.0001	0.333	0.296	0.375
Skin Infection and Inflammation (197, 198, 200)	30,972	1.68	-0.962	0.016	<.0001	0.382	0.370	0.394
Chronic Ulcer of Skin (199)	6,013	0.33	-0.864	0.032	<.0001	0.421	0.395	0.449
Spondylosis, Intervertebral Disc Disorders, Other Back Problems (205)	28,209	1.53	-0.604	0.016	<.0001	0.547	0.530	0.565
Pathological Fracture (207)	13,499	0.73	-0.950	0.020	<.0001	0.387	0.372	0.402
Rheumatoid Arthritis and Other Connective Tissue Disease (202, 210, 211)	15,813	0.86	-0.716	0.020	<.0001	0.489	0.470	0.508
Acquired Deformities and Congenital Anomalies (208, 209, 212, 213, 214, 215, 216, 217)	10,822	0.59	-0.600	0.024	<.0001	0.549	0.524	0.576
Joint Disorders, Dislocation, and Fractures (225, 228, 229, 230, 231)	96,584	5.23	-0.816	0.011	<.0001	0.442	0.432	0.452
Fracture of Neck of Femur (Hip) (226)	117,028	6.34	-1.067	0.011	<.0001	0.344	0.337	0.351
Spinal Cord Injury (227)	842	0.05	-1.306	0.074	<.0001	0.271	0.234	0.314
Sprains and Strains (232)	2,515	0.14	-0.411	0.046	<.0001	0.663	0.606	0.725

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled
Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
Intracranial Injury (233)	17,808	0.96	-1.044	0.020	<.0001	0.352	0.339	0.366
Crushing Injury or Internal Injury (234)	3,574	0.19	-0.710	0.037	<.0001	0.492	0.457	0.529
Open Wounds and Burns (235, 236, 240)	2,864	0.16	-0.913	0.041	<.0001	0.401	0.370	0.435
Complications of Device, Procedures, or Medical Care (237, 238)	83,520	4.52	-0.947	0.012	<.0001	0.388	0.379	0.397
Superficial Injury, Contusion (239)	4,873	0.26	-0.748	0.032	<.0001	0.473	0.445	0.504
Poisoning and Injury (241, 242, 243, 260)	3,597	0.19	-0.607	0.037	<.0001	0.545	0.507	0.586
Other Injuries and Conditions due to External Causes (244)	3,768	0.20	-1.124	0.037	<.0001	0.325	0.302	0.349
Syncope (245)	9,704	0.53	-0.759	0.024	<.0001	0.468	0.447	0.490
Fever of Unknown Origin (246)	1,752	0.09	-1.130	0.053	<.0001	0.323	0.291	0.359
Gangrene (248)	5,628	0.30	-1.459	0.034	<.0001	0.233	0.218	0.248
Shock (249)	167	0.01	-1.100	0.173	<.0001	0.333	0.237	0.467
Nausea and Vomiting (250)	1,226	0.07	-1.066	0.062	<.0001	0.344	0.305	0.389
Abdominal Pain (251)	2,019	0.11	-1.028	0.049	<.0001	0.358	0.325	0.394
Malaise and Fatigue (252)	4,222	0.23	-0.925	0.034	<.0001	0.397	0.371	0.424
Allergic Reactions (253)	527	0.03	-0.838	0.093	<.0001	0.433	0.360	0.519
Rehabilitation Care, Fitting of Prostheses, and Adjustment of Devices (254)	1,561	0.08	-0.141	0.056	0.0119	0.868	0.778	0.969
Other Aftercare (257)	729	0.04	-0.626	0.083	<.0001	0.535	0.454	0.629
Residual Codes, Unclassified (259)	8,638	0.47	-1.142	0.026	<.0001	0.319	0.303	0.336
Psychosocial Disorders (650, 651, 652, 654, 655, 656)	1,045	0.06	-1.307	0.076	<.0001	0.271	0.233	0.314
Delirium (653)	30,350	1.64	-1.565	0.021	<.0001	0.209	0.201	0.218
Mood and Personality Disorders (657, 658, 662)	10,572	0.57	-1.037	0.031	<.0001	0.355	0.334	0.377
Schizophrenia and Other Psychotic Disorders (659)	10,713	0.58	-1.641	0.032	<.0001	0.194	0.182	0.206

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled
Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
Substance Abuse (660, 661, 663)	8,046	0.44	-0.741	0.026	<.0001	0.477	0.453	0.502
Miscellaneous Disorders (670)	293	0.02	-0.666	0.125	<.0001	0.514	0.402	0.657
Surgical Categories based on Prior Acute Stay								
Vascular	15,019	0.81	0.234	0.021	<.0001	1.264	1.213	1.317
Orthopedic	363,399	19.69	0.316	0.008	<.0001	1.371	1.350	1.392
General	92,519	5.01	0.314	0.010	<.0001	1.369	1.343	1.396
Cardio Thoracic	36,348	1.97	0.639	0.016	<.0001	1.895	1.837	1.955
Urologic	16,079	0.87	0.082	0.021	<.0001	1.086	1.043	1.131
Neurosurgery	15,363	0.83	0.120	0.020	<.0001	1.127	1.085	1.171
Plastic	27,846	1.51	0.074	0.014	<.0001	1.076	1.047	1.107
Otolaryngology	3,372	0.18	0.194	0.041	<.0001	1.215	1.122	1.315
Obstetrics/Gynecology	4,376	0.24	0.313	0.038	<.0001	1.367	1.269	1.474
Dialysis in Prior Acute Stay where End-Stage Renal Disease not Indicated							ı	
Dialysis where End-Stage Renal Disease Not Indicated	9,898	0.54	-0.069	0.024	0.0032	0.933	0.891	0.977
End-Stage Renal Disease								
End-Stage Renal Disease Indicator at the Time of Index SNF	83,674	4.53	-0.447	0.017	<.0001		0.618	0.661
Prior Acute Length of Stay in Non-Psychiatric Hospital or Prior Stay in Psy		<b>`</b>	1			1	ı	
Prior Acute Stay in Psychiatric Hospital	35,233	1.91	-0.883	0.022	<.0001	0.414	0.396	0.432
4-7 Days in Non-Psychiatric Hospital	823,517	44.61	-0.079	0.004	<.0001	0.924	0.916	0.932
8-14 Days in Non-Psychiatric Hospital	370,664	20.08	-0.220	0.005	<.0001	0.802	0.794	0.811
15-29 Days in Non-Psychiatric Hospital	111,630	6.05	-0.401	0.008	<.0001	0.669	0.658	0.681
>29 Days in Non-Psychiatric Hospital	16,914	0.92	-0.681	0.020	<.0001	0.506	0.486	0.526
Ventilator Use in SNF								
Ventilator Use During Index SNF Stay (MedPAR Diag Code = Respirator (V4611))	2,208	0.12	-1.255	0.096	<.0001	0.285	0.236	0.344

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
Comorbidities - Hierarchical Condition Categories (HCCs) (* indicates that based on acute care claims from the past 365 days (including the most rece			ne most recent	acute c	are claim.	HCCs not pr	eceded by * a	ire
HCC1: HIV/AIDS	5,048	0.27	-0.173	0.034	<.0001	0.841	0.787	0.899
HCC2: Septicemia, Sepsis, Systemic Inflammatory Response Syndrome/Shock*	197,021	10.67	-0.090	0.010	<.0001	0.914	0.897	0.932
HCC6: Opportunistic Infections	18,132	0.98	-0.030	0.018	0.0918	0.971	0.938	1.005
HCC7: Other Infectious Diseases*	310,312	16.81	-0.037	0.005	<.0001	0.964	0.955	0.973
HCC8: Metastatic Cancer and Acute Leukemia	48,788	2.64	-0.739	0.011	<.0001	0.478	0.467	0.488
HCC9: Lung and Other Severe Cancers	31,994	1.73	-0.371	0.013	<.0001	0.690	0.673	0.708
HCC10: Lymphoma and Other Cancers	28,997	1.57	-0.153	0.013	<.0001	0.858	0.836	0.881
HCC11: Colorectal, Bladder, and Other Cancers	21,538	1.17	-0.066	0.015	<.0001	0.936	0.909	0.965
HCC17-HCC19: Diabetes with Acute Complications; Diabetes with Chronic Complications; Diabetes without Complication	695,120	37.65	-0.125	0.004	<.0001	0.882	0.876	0.889
HCC20: Type I Diabetes Mellitus	20,238	1.10	-0.076	0.017	<.0001	0.927	0.897	0.959
HCC21: Protein-Calorie Malnutrition	277,664	15.04	-0.239	0.005	<.0001	0.787	0.779	0.795
HCC23: Other Significant Endocrine and Metabolic Disorders	113,500	6.15	-0.007	0.007	0.3165	0.993	0.978	1.007
HCC24: Disorders of Fluid/Electrolyte/Acid-Base Balance	1,051,719	56.97	-0.065	0.004	<.0001	0.937	0.930	0.944
HCC27; HCC28: End-stage liver disease; Cirrhosis of Liver	47,723	2.59	-0.343	0.012	<.0001	0.709	0.693	0.726
HCC29: Chronic Hepatitis	9,562	0.52	-0.063	0.024	0.0088	0.939	0.896	0.984
HCC36: Peptic Ulcer, Hemorrhage, Other Specified Gastrointestinal Disorders*	128,566	6.96	-0.080	0.007	<.0001	0.923	0.911	0.936
HCC40: Rheumatoid Arthritis and Inflammatory Connective Tissue Disease	111,971	6.07	-0.011	0.007	0.1226	0.989	0.976	1.003
HCC46: Severe Hematological Disorders	21,649	1.17	-0.229	0.016	<.0001	0.795	0.770	0.821
HCC47: Disorders of Immunity	55,481	3.01	-0.058	0.011	<.0001	0.944	0.925	0.964

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
HCC48: Coagulation Defects and Other Specified Hematological Disorders	214,702	11.63	-0.011	0.006	0.0624	0.989	0.978	1.001
HCC49: Iron Deficiency and Other/Unspecified Anemias and Blood Disease	834,200	45.19	-0.063	0.004	<.0001	0.939	0.932	0.946
HCC51; HCC52: Dementia With Complications; Dementia Without Complication	571,733	30.97	-0.753	0.004	<.0001	0.471	0.467	0.475
HCC53: Nonpsychotic Organic Brain Syndromes/Conditions	19,207	1.04	-0.083	0.016	<.0001	0.920	0.893	0.949
HCC57: Schizophrenia	43,724	2.37	-0.837	0.013	<.0001	0.433	0.422	0.444
HCC58: Major Depressive, Bipolar, and Paranoid Disorders	94,242	5.11	-0.189	0.008	<.0001	0.828	0.815	0.841
HCC59: Reactive and Unspecified Psychosis	44,748	2.42	-0.202	0.011	<.0001	0.817	0.800	0.835
HCC60: Personality Disorders	3,444	0.19	-0.350	0.040	<.0001	0.705	0.651	0.763
HCC61: Depression	368,707	19.97	-0.123	0.004	<.0001	0.885	0.877	0.892
HCC63: Other Psychiatric Disorders	125,636	6.81	-0.053	0.007	<.0001	0.948	0.936	0.960
HCC64-HCC66: Profound Mental Retardation/Developmental Disability; Severe Mental Retardation/Developmental Disability; Moderate Mental Retardation/Developmental Disability	3,952	0.21	-0.612	0.041	<.0001	0.542	0.501	0.587
HCC67; HCC68: Mild Mental Retardation, Autism, Down's Syndrome: Other Developmental Disability	16,866	0.91	-0.548	0.019	<.0001	0.578	0.557	0.600
HCC70: Quadriplegia	11,227	0.61	-0.730	0.026	<.0001	0.482	0.458	0.507
HCC71: Paraplegia	12,106	0.66	-0.371	0.022	<.0001	0.690	0.660	0.721
HCC72: Spinal Cord Disorders/Injuries	10,404	0.56	-0.074	0.022	0.0008	0.929	0.889	0.970
HCC73: Amyotrophic Lateral Sclerosis and Other Motor Neuron Disease	1,218	0.07	-0.453	0.068	<.0001	0.636	0.556	0.727
HCC74: Cerebral Palsy	5,575	0.30	-0.419	0.033	<.0001	0.658	0.617	0.701
HCC76: Muscular Dystrophy	1,437	0.08	-0.322	0.061	<.0001	0.725	0.644	0.816

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled
Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
HCC77: Multiple Sclerosis	13,228	0.72	-0.316	0.021	<.0001	0.729	0.700	0.759
HCC78: Parkinson's and Huntington's Diseases	71,304	3.86	-0.167	0.009	<.0001	0.847	0.832	0.861
HCC79: Seizure Disorders and Convulsions	126,557	6.86	-0.167	0.007	<.0001	0.846	0.835	0.858
HCC80: Coma, Brain Compression/Anoxic Damage	26,262	1.42	-0.208	0.016	<.0001	0.813	0.788	0.838
HCC82: Respirator Dependence/Tracheostomy Status	16,905	0.92	-0.338	0.022	<.0001	0.713	0.683	0.745
HCC84: Cardio-Respiratory Failure and Shock	342,239	18.54	-0.063	0.005	<.0001	0.939	0.930	0.949
HCC85: Congestive Heart Failure	708,821	38.40	-0.180	0.004	<.0001	0.835	0.829	0.842
HCC86: Acute Myocardial Infarction	106,088	5.75	-0.040	0.007	<.0001	0.961	0.947	0.975
HCC90: Heart Infection/Inflammation, Except Rheumatic*	12,254	0.66	-0.011	0.020	0.5765	0.989	0.950	1.029
HCC96: Specified Heart Arrhythmias	646,780	35.04	-0.070	0.004	<.0001	0.932	0.926	0.939
HCC99: Cerebral Hemorrhage*	7,278	0.39	-0.166	0.027	<.0001	0.847	0.804	0.893
HCC100: Ischemic or Unspecified Stroke*	15,400	0.83	-0.172	0.018	<.0001	0.842	0.812	0.873
HCC103: Hemiplegia/Hemiparesis	110,982	6.01	-0.439	0.008	<.0001	0.645	0.635	0.655
HCC104: Monoplegia, Other Paralytic Syndromes	4,698	0.25	-0.195	0.033	<.0001	0.822	0.771	0.877
HCC105: Late Effects of Cerebrovascular Disease, Except Paralysis	81,880	4.44	-0.083	0.008	<.0001	0.921	0.906	0.936
HCC106: Atherosclerosis of the Extremities with Ulceration or Gangrene	41,309	2.24	-0.324	0.013	<.0001	0.723	0.705	0.742
HCC107; HCC108: Vascular Disease with Complications; Vascular Disease*	232,547	12.60	-0.023	0.005	<.0001	0.977	0.968	0.987
HCC110; HCC111: Cystic Fibrosis; Chronic Obstructive Pulmonary Disease	547,331	29.65	-0.149	0.004	<.0001	0.861	0.855	0.868

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled
Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
HCC114: Aspiration and Specified Bacterial Pneumonias	135,458	7.34	-0.244	0.007	<.0001	0.784	0.772	0.795
HCC116: Viral and Unspecified Pneumonia, Pleurisy	303,351	16.43	-0.144	0.005	<.0001	0.866	0.858	0.875
HCC117: Pleural Effusion/Pneumothorax	135,479	7.34	-0.048	0.007	<.0001	0.953	0.941	0.966
HCC119: Legally Blind	28,492	1.54	-0.158	0.014	<.0001	0.854	0.831	0.877
HCC120: Major Eye Infections/Inflammations	1,563	0.08	-0.157	0.059	0.0077	0.855	0.761	0.959
HCC132: Kidney Transplant Status	8,535	0.46	-0.183	0.028	<.0001	0.833	0.789	0.880
HCC134: Dialysis Status	63,441	3.44	-0.351	0.019	<.0001	0.704	0.678	0.731
HCC135: Acute Renal Failure	531,060	28.77	-0.151	0.004	<.0001	0.860	0.852	0.867
HCC136: Chronic Kidney Disease (Stage 5)	9,504	0.51	-0.394	0.026	<.0001	0.674	0.640	0.710
HCC137: Chronic Kidney Disease, Severe (Stage 4)	20,295	1.10	-0.267	0.016	<.0001	0.766	0.742	0.790
HCC138; HCC139: Chronic Kidney Disease, Moderate (Stage 3); Chronic Kidney Disease, Mild or Unspecified (Stages 1-2 or Unspecified)	173,051	9.37	-0.108	0.006	<.0001	0.898	0.888	0.908
HCC140: Unspecified Renal Failure	1,201	0.07	-0.371	0.066	<.0001	0.690	0.607	0.785
HCC144: Urinary Tract Infection	600,227	32.51	-0.212	0.004	<.0001	0.809	0.802	0.815
HCC145: Other Urinary Tract Disorders	208,407	11.29	-0.023	0.005	<.0001	0.977	0.967	0.988
HCC157: Pressure Ulcer of Skin with Necrosis Through to Muscle, Tendon, or Bone	17,694	0.96	-0.757	0.023	<.0001	0.469	0.448	0.490
HCC158: Pressure Ulcer of Skin with Full Thickness Skin Loss	37,560	2.03	-0.619	0.014	<.0001	0.539	0.524	0.554
HCC159: Pressure Ulcer of Skin with Partial Thickness Skin Loss	55,374	3.00	-0.443	0.011	<.0001	0.642	0.628	0.656
HCC160: Pressure Pre-Ulcer Skin Changes or Unspecified Stage	47,314	2.56	-0.393	0.011	<.0001	0.675	0.660	0.690
HCC161: Chronic Ulcer of Skin, Except Pressure	61,215	3.32	-0.134	0.010	<.0001	0.874	0.858	0.891
HCC164: Cellulitis, Local Skin Infection	182,826	9.90	-0.036	0.006	<.0001	0.965	0.953	0.976
HCC166; HCC167: Severe Head Injury; Major Head Injury*	10,594	0.57	-0.091	0.022	<.0001	0.913	0.875	0.953

Table 1-1. Preliminary Logistic Regression Model Results for Discharge to Community-Post Acute Care (PAC) Skilled Nursing Facility Quality Reporting Program, 2013 (continued)

Covariate	N	%	Coefficient Estimate	SE <sup>1</sup>	p- value	Odds Ratio (OR)	OR 95% Lower CL <sup>2</sup>	OR 95% Upper CL
HCC176: Complications of Specified Implanted Device or Graft	88,079	4.77	-0.041	0.009	<.0001	0.960	0.944	0.976
HCC188: Artificial Openings for Feeding or Elimination	68,440	3.71	-0.254	0.010	<.0001	0.775	0.760	0.791
HCC189; HCC190: Amputation Status, Lower Limb/Amputation Complications; Amputation Status, Upper Limb		1.86	-0.244	0.013	<.0001	0.784	0.764	0.804
Acute History: Number of Hospital Stays in Past Year, Excluding Most Rec	ent Stay (Re	ference: 0-3	3 Stays)		•	1		
4-6 Stays	141,879	7.69	-0.097	0.007	<.0001	0.907	0.895	0.920
7-9 Stays	26,690	1.45	-0.296	0.017	<.0001	0.744	0.720	0.769
>10 Stays	8,782	0.48	-0.700	0.033	<.0001	0.497	0.466	0.530

<sup>&</sup>lt;sup>1</sup> SE = Standard Error; <sup>2</sup> CL = Confidence Limit.

Source: RTI International analysis of Medicare claims data (RTI program reference: Model 13C).

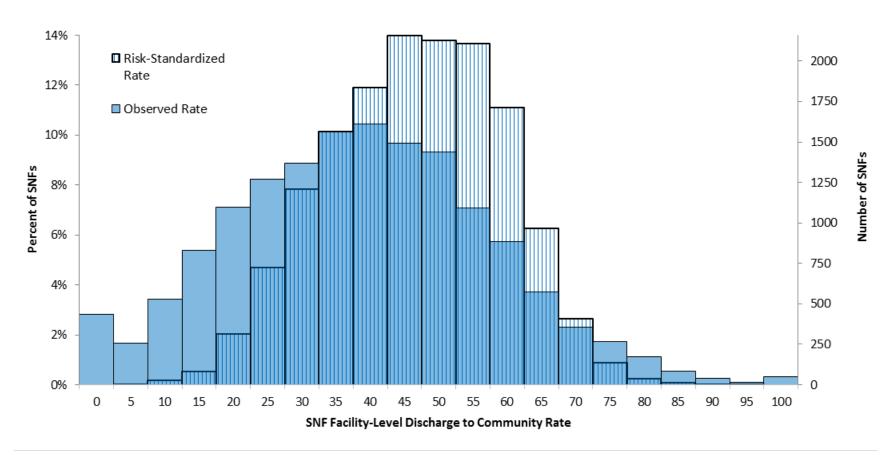
Table 1-2. Skilled Nursing Facility: Facility-Level Observed and Risk-Standardized Discharge to Community Rates, 2013

Discharge to Community Rate	Mean	SD	Min	1st pctl	5th petl	10th pctl	25th pctl	50th pctl (Median)	75th petl	90th pctl	95th pctl	99th pctl	Max
Observed	36.49	18.59	0.00	0.00	6.06	12.50	23.08	36.09	49.09	60.16	67.68	81.82	100.00
Risk-Standardized	44.09	12.74	3.07	15.98	22.82	26.80	34.82	44.48	53.63	60.10	63.84	71.11	94.81

NOTE: Based on CY 2013 Medicare fee-for-service claims data from 15,430 SNFs. Facility-level number of SNF stays ranged from 1 to 1,683, with a mean of 119.64 and median of 83. SD = standard deviation, pctl = percentile.

Source: RTI analysis of Medicare claims data (RTI program reference: Model 13C).

Figure 1-1. Skilled Nursing Facility: Facility-Level Observed and Risk-Standardized Discharge to Community Rates, 2013



Note: Based on CY 2013 Medicare fee-for-service claims data from 15,430 SNFs. Facility-level number of SNF stays ranged from 1 to 1,683, with a mean of 119.64 and median of 83. Blue bars represent the observed rate distribution; striped bars represent the risk-standardized rate distribution; the overlap between blue and striped bars represents the overlap between observed and risk-standardized rate distributions.

Source: RTI analysis of Medicare claims data (RTI program reference: Model 13C).

#### **APPENDIX 2**

# POTENTIALLY PREVENTABLE 30-DAY POST-DISCHARGE READMISSION MEASURE FOR SKILLED NURSING FACILITY (SNF) QUALITY REPORTING PROGRAM (QRP)

- o Table 2-1. List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-9 Codes
- Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes
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Table 2-1. List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-9 Codes

Conditions	These conditions will be used Diagnosis	ICD-9- CM	30 day post- PAC discharge	Clinical Rationale
Adult asthma*	*Extrinsic asthma NOS	493.00	X	Inadequate management of chronic conditions
	*Ext asthma w/ status asth	493.01	X	
	*Ext asthma w(acute) exac	493.02	X	
	*Intrinsic asthma NOS	493.10	X	
	*Int asthma w status asth	493.11	X	
	*Int asthma w (ac) exac	493.12	X	
	*Chronic obst asthma NOS	493.20	X	
	*Ch ob asthma w stat asth	493.21	X	
	*Ch obst asth w (ac) exac	493.22	X	
	*Exercise ind bronchospasm	493.81	X	
	*Cough variant asthma	493.82	X	
	*Asthma NOS	493.90	X	
	*Asthma w status asth mat	493.91	X	
	*Asthma NOS w (ac) exac	493.92	X	
Chronic obstructive pulmonary disease (COPD)*	*Simple Chr Bronchitis	491.0	X	Inadequate management of chronic conditions
	*Mucopurul Chr Bronchitis	491.1	X	
	*Obs Chr Brnc w/o act exa	491.20	X	
	*Obs Chr Brnc w/ act exa	491.21	X	
	*Obs Chr Bronc w/ ac Bronc	491.22	X	
	*Chronic Bronchitis NEC	491.8	X	
	*Chronic Bronchitis NOS	491.9	X	
	*Emphysematous Bleb	492.0	X	
	*Emphysema NEC	492.8	X	
	*Bronchiectasis	494	X	
	*Bronchiectas w/o ac exac	494.0	X	
	*Bronchiectasis w/ ac exac	494.1	X	
	*Chr airway obstruct NEC	496	X	

Table 2-1. List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-9 Codes (continued)

Conditions	Diagnosis	ICD-9- CM	30 day post- PAC discharge	Clinical Rationale
Congestive heart	*Rheumatic Heart Failure	398.91	X	Inadequate
failure (CHF)*	Tricumatic Treat Fantice	350.51		management of
				chronic conditions
	*Mal hypert hrt dis w/ CHF	402.01	X	
	*Benign hyp hrt dis w CHF	402.11	X	
	*Hyperten heart dis w CHF	402.91	X	
	*Mal hyper hrt/ren w/ CHF	404.01	X	
	*Mal hyp hrt/ren w CHF/RF	404.03	X	
	*Ben hyper hrt/ren w CHF	404.11	X	
	*Ben hyp hrt/ren w CHF/RF	404.13	X	
	*Hyper hrt/ren NOS w CHF	404.91	X	
	*Hyp Ht/Ren NOS w CHR	404.93	X	
	*Congestive Heart Failure	428.0	X	
	*Left heart failure	428.1	X	
	*Systolic hrt failure NOS	428.20	X	
	*AC systolic hrt failure	428.21	X	
	*Chr systolic hrt failure	428.22	X	
	*AC on chr syst hrt fail	428.23	X	
	*Diastolic hrt failure NOS	428.30	X	
	*AC diastolic hrt failure	428.31	X	
	*Chr diastolic hrt fail	428.32	X	
	*AC on chr diast hrt fail	428.33	X	
	*Syst/diast hrt fail NOS	428.40	X	
	*AC syst/diastole hrt fail	428.41	X	
	*Chr syst/diastl hrt fail	428.42	X	
	*AC/CHR syst/dia hrt fail	428.43	X	
	*Heart Failure NOS	428.9	X	
	Acute lung edema NOS	518.4	X	1
Diabetes short-term complication*	Secondary diabetes mellitus with ketoacidosis	249.1X	X	Inadequate management of
Complication	Recoderdosis			chronic conditions
	Secondary diabetes mellitus with hyperosmolarity	249.2X	X	
	Secondary diabetes mellitus with other coma	249.3X	X	

Table 2-1. List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-9 Codes (continued)

Conditions	Diagnosis	ICD-9- CM	30 day post- PAC discharge	Clinical Rationale
	Secondary diabetes mellitus with other specified manifestations (hypoglycemia)	249.8X	X	
	Diabetes with other specified manifestations (hypoglycemia)	250.8X	X	
	*DM Keto T2, DM Cont	250.10	X	
	*DM Keto T1, DM Cont	250.11	X	
	*DM Keto T2, DM Uncont	250.12	X	
	*DM Keto T1, DM Uncont	250.13	X	
	*DM W/ Hyprosm T2, DM Cont	250.20	X	
	*DM W/ Hyprosm T1, DM Cont	250.21	X	
	*DM W/ Hyprosm T2, DM Uncnt	250.22	X	
	*DM W/ Hyprosm T1, DM Uncnt	250.23	X	
	*DM Coma Nec Typ Ii, DM Cnt	250.30	X	
	*DM Coma Nec T1, DM Cont	250.31	X	
	*DM Coma Nec T2, DM Uncont	250.32	X	
	*DM Coma Nec T1, DM Uncont	250.33	X	
	*Malignant Hypertension	401.0	X	Inadequate management of chronic conditions
	*Hypertension NOS	401.9	X	
	*Mal Hyperten hrt dis NOS	402.00	X	
	*Benign hyp ht dis w/o hf	402.10	X	
	*Hyp hrt dis NOS w/o hf	402.90	X	
	*Mal hyp ren w/o ren fail	403.00	X	
	*Ben hy kid w cr kid I-IV	403.10	X	
	*Hy kid NOS w cr kid I-IV	403.90	X	
	*Mal hy ht/ren w/o chf/rf	404.00	X	
	*Ben hy ht/ren w/o chf/rf	404.10	X	
	*Hy ht/ren NOS w/o chf/rf	404.90	X	
	Orthostatic hypotension	458.0	X	
	Chronic hypotension	458.1	X	
	Iatrogenic hypotension NEC	458.29	X	
	Hypotension NEC	458.8	X	
	Hypotension NOS	458.9	X	

Table 2-1. List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-9 Codes (continued)

Conditions	Diagnosis	ICD-9- CM	30 day post- PAC discharge	Clinical Rationale
Influenza	Influenza	487.X	X	Inadequate
	Influenza due to identified avian influenza virus	488.X	X	management of infection
Bacterial pneumonia*	*Pneumococcal Pneumonia	481	X	Inadequate management of infection
	*H.Influenzae Pneumonia	482.2	X	
	*Strep Pneumonia Unspec	482.30	X	
	*Grp A Strep Pneumonia	482.31	X	
	*Grp B Strep Pneumonia	482.32	X	
	*Oth Strep Pneumonia	482.39	X	
	*Meth Sus Pneum D/T Staph	482.41	X	
	*Meth Res Pneu D/T Staph	482.42	X	
	*Bacterial Pneumonia Nos	482.9	X	
	*Mycoplasma Pneumonia	483.0	X	
	*Chlamydia Pneumonia	483.1	X	
	*Oth Spec Org Pneumonia	483.8	X	
	*Broncopneumonia Org Nos	485	X	
	*Pneumonia, Organism Nos	486	X	
Urinary tract infection*/Kidney infection	*Ac pyelonephritis NOS	590.10	X	Inadequate management of infection
	*Ac pyelonephr w med necr	590.11	X	
	*Renal/perirenal abscess	590.2	X	
	*Pyeloureteritis cystica	590.3	X	
	*Pyelonephritis NOS	590.80	X	
	*Pyelonephrit in oth dis	590.81	X	
	*Infection of kidney NOS	590.9	X	
	*Acute cystitis	595.0	X	
	Urethral abscess	597.0	X	
	*Urin tract infection NOS	599.0	X	
C. difficile infection [135 subset]	Intestinal infection due to Clostridium difficile	008.45	X	Inadequate management of infection

Table 2-1. List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-9 Codes (continued)

	nese conditions will be used f		30 day post-	
Conditions	Diagnosis	ICD-9- CM	PAC discharge	Clinical Rationale
Septicemia (except in labor) [2]	Salmonella septicemia	003.1	X	Inadequate management of infection
	Septicemic plague	020.2	X	
	Anthrax septicemia	022.3	X	
	Meningococcemia	036.2	X	
	Streptococcal septicemia	038.0	X	
	Staphylococcal septicemia	038.1	X	
	Staphylococcal septicemia, unspecified	038.10	X	
	Methicillin susceptible Staphylococcus aureus septicemia	038.11	X	
	Methicillin resistant Staphylococcus aureus septicemia	038.12	X	
	Other staphylococcal septicemia	038.19	X	
	Pneumococcal septicemia [Streptococcus pneumoniae septicemia]	038.2	X	
	Septicemia due to anaerobes	038.3	X	
	Septicemia due to gram-negative organism, unspecified	038.40	X	
	Septicemia due to hemophilus influenzae [H. influenzae]	038.41	X	
	Septicemia due to escherichia coli [E. coli]	038.42	X	
	Septicemia due to pseudomonas	038.43	X	
	Septicemia due to serratia	038.44	X	
	Other septicemia due to gram- negative organisms	038.49	X	
	Other specified septicemias	038.8	X	
	Unspecified septicemia	038.9	X	
	Herpetic septicemia	054.5	X	
	Septic arterial embolism	449	X	
	Sepsis	995.91	X	
	Severe sepsis	995.92	X	
	Septic shock	785.52	X	

Table 2-1. List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-9 Codes (continued)

1,010.	these conditions will be used f	or the post 1	30 day post-	
Conditions	Diagnosis	ICD-9- CM	PAC discharge	Clinical Rationale
Skin and subcutaneous tissue infections [197]	Cellulitis and abscess of finger, unspecified	681.00	X	Inadequate management of infection
	Cellulitis and abscess of toe, unspecified	681.10	X	
	Cellulitis and abscess of unspecified digit	681.9	X	
	Cellulitis and abscess of face	682.0	X	
	Cellulitis and abscess of neck	682.1	X	
	Cellulitis and abscess of trunk	682.2	X	
	Cellulitis and abscess of upper arm and forearm	682.3	X	
	Cellulitis and abscess of hand, except fingers and thumb	682.4	X	
	Cellulitis and abscess of buttock	682.5	X	
	Cellulitis and abscess of leg, except foot	682.6	X	
	Cellulitis and abscess of foot, except toes	682.7	X	
	Cellulitis and abscess of other specified sites	682.8	X	
	Cellulitis and abscess of unspecified sites	682.9	X	
	Other specified local infections of skin and subcutaneous tissue	686.8	X	
	Unspecified local infection of skin and subcutaneous tissue	686.9	X	
Dehydration*/ Electrolyte imbalance [55]	**Hyperosmolality and/or hypernatremia	276.0	X	Inadequate management of other unplanned events
	Hyposmolality and/or hyponatremia	276.1	X	
	Acidosis	276.2	X	
	Alkalosis	276.3	X	]
	Mixed acid-base balance disorder	276.4	X	]
	*Volume depletion, unspecified	276.50	X	]
	*Dehydration	276.51	X	

Table 2-1. List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-9 Codes (continued)

1,010.	These conditions will be used f 	or the post 1	30 day post-	ecusures.
Conditions	Diagnosis	ICD-9- CM	PAC discharge	Clinical Rationale
	*Hypovolemia	276.52	X	
	Fluid overload disorder	276.6	X	
	Other fluid overload	276.69	X	
	Hyperpotassemia	276.7	X	
	Hypopotassemia	276.8	X	
	Electrolyte and fluid disorders not elsewhere classified	276.9	X	
	**Intes Infec Rotavirus	008.61	X	
	**Intes Infec Adenovirus	008.62	X	
	**Int Inf Norwalk Virus	008.63	X	
	**Int Inf Oth Sml Rnd Vrus	008.64	X	
	**Intes Infec Calcivirus	008.65	X	
	**Intes Infec Astrovirus	008.66	X	
	**Int Inf Enterovirus NEC	008.67	X	
	**Enteritis NOS	008.69	X	
	**Viral Enteritis NOS	008.8	X	
	**Infectious Enteritis NOS	009.0	X	
	**Enteritis of Infect Orig	009.1	X	
	**Infectious Diarrhea NOS	009.2	X	
	**Diarrhea of Infect Orig	009.3	X	
	**Noninf Gastroenterit NEC	558.9	X	
Aspiration pneumonitis; food/vomitus [129]	Pneumonitis due to inhalation of food or vomitus	507.0	X	Inadequate management of other unplanned events
Acute renal failure*	*Acute kidney failure with lesion of tubular necrosis	584.5	X	Inadequate management of other unplanned events
	*Acute kidney failure with lesion of renal cortical necrosis	584.6	X	
	*Acute kidney failure with lesion of renal medullary [papillary] necrosis	584.7	X	
	*Acute kidney failure with other specified pathological lesion in kidney	584.8	X	
	*Acute kidney failure, unspecified	584.9	X	
	*Renal Failure NOS	586	X	

Table 2-1. List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-9 Codes (continued)

Conditions	Diagnosis	ICD-9- CM	30 day post- PAC discharge	Clinical Rationale
	*Surg Compl-Urinary Tract	997.5	X	
Arrhythmia	Atrial fibrillation	427.31	X	Inadequate management of other unplanned events
	Atrial flutter	427.32	X	
Intestinal impaction [145 subset]	Impaction of intestine, unspecified	560.30	X	Inadequate management of other unplanned events
	Fecal impaction	560.32	X	
	Other impaction of intestine	560.39	X	
Pressure ulcers	Chronic ulcer of skin	707.0X 707.2X	X	Inadequate management of other unplanned events

SOURCE: List of potentially preventable readmission conditions from RTI International with ICD-9-CM (version: April 2016).

NOTES: [###] indicates Clinical Classifications Software (CCS) code

To be considered a potentially preventable readmission, diagnosis codes must be the principal diagnosis on the readmission claim, except where noted.

<sup>\*</sup>Ambulatory Care Sensitive Conditions (ACSCs)/Prevention Quality Indicators (PQIs)

<sup>\*\*</sup> Primary diagnosis with dehydration (codes: 276.50, 276.51, 276.52) as secondary diagnosis

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
Adult Asthma*	Asthma* (PQI 05)	J4521	Mild intermittent asthma with (acute) exacerbation
		J4522	Mild intermittent asthma with status asthmaticus
		J4531	Mild persistent asthma with (acute) exacerbation
		J4532	Mild persistent asthma with status asthmaticus
		J4541	Moderate persistent asthma with (acute) exacerbation
		J4542	Moderate persistent asthma with status asthmaticus
		J4551	Severe persistent asthma with (acute) exacerbation
		J4552	Severe persistent asthma with status asthmaticus
		J45901	Unspecified asthma with (acute) exacerbation
		J45902	Unspecified asthma with status asthmaticus
		J45990	Exercise induced bronchospasm
		J45991	Cough variant asthma
		J45998	Other asthma
	Acute Bronchitis*^ (PQI 05)	J200	Acute bronchitis due to Mycoplasma pneumoniae
		J201	Acute bronchitis due to Hemophilus influenzae
		J202	Acute bronchitis due to streptococcus
		J203	Acute bronchitis due to coxsackievirus
		J204	Acute bronchitis due to parainfluenza virus
		J205	Acute bronchitis due to respiratory syncytial virus
		J206	Acute bronchitis due to rhinovirus
		J207	Acute bronchitis due to echovirus
		J208	Acute bronchitis due to other specified organisms
		J209	Acute bronchitis, unspecified
		J40	Bronchitis, not specified as acute or chronic
Chronic obstructive	COPD* (PQI 05)	J410	Simple chronic bronchitis
pulmonary disease (COPD)		J411	Mucopurulent chronic bronchitis
(COLD)		J418	Mixed simple and mucopurulent chronic bronchitis

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		J42	Unspecified chronic bronchitis
		J430	Unilateral pulmonary emphysema [MacLeod's syndrome]
		J431	Panlobular emphysema
		J432	Centrilobular emphysema
		J438	Other emphysema
		J439	Emphysema, unspecified
		J440	Chronic obstructive pulmonary disease with acute lower respiratory infection
		J441	Chronic obstructive pulmonary disease with (acute) exacerbation
		J449	Chronic obstructive pulmonary disease, unspecified
		J470	Bronchiectasis with acute lower respiratory infection
		J471	Bronchiectasis with (acute) exacerbation
		J479	Bronchiectasis, uncomplicated
Congestive heart		I09.81	Rheumatic heart failure
failure (CHF)		I11.0	Hypertensive heart disease with heart failure
		I11.0	Hypertensive heart disease with heart failure
		I11.0	Hypertensive heart disease with heart failure
		113.0	Hypertensive heart and chronic kidney disease with heart failure and stage 1 through stage 4 chronic kidney disease, or unspecified chronic kidney disease
		I13.2	Hypertensive heart and chronic kidney disease with heart failure and with stage 5 chronic kidney disease, or end stage renal disease
		I13.0	Hypertensive heart and chronic kidney disease with heart failure and stage 1 through stage 4 chronic kidney disease, or unspecified chronic kidney disease
		I13.2	Hypertensive heart and chronic kidney disease with heart failure and with stage 5 chronic kidney disease, or end stage renal disease
		113.0	Hypertensive heart and chronic kidney disease with heart failure and stage 1 through stage 4 chronic kidney disease, or unspecified chronic kidney disease

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		I13.2	Hypertensive heart and chronic kidney disease with heart failure and with stage 5 chronic kidney disease, or end stage renal disease
		I50.9	Heart failure, unspecified
		I50.1	Left ventricular failure
		I50.20	Unspecified systolic (congestive) heart failure
		I50.21	Acute systolic (congestive) heart failure
		I50.22	Chronic systolic (congestive) heart failure
		150.23	Acute on chronic systolic (congestive) heart failure
		150.30	Unspecified diastolic (congestive) heart failure
		I50.31	Acute diastolic (congestive) heart failure
		150.32	Chronic diastolic (congestive) heart failure
		150.33	Acute on chronic diastolic (congestive) heart failure
		150.40	Unspecified combined systolic (congestive) and diastolic (congestive) heart failure
		I50.41	Acute combined systolic (congestive) and diastolic (congestive) heart failure
		150.42	Chronic combined systolic (congestive) and diastolic (congestive) heart failure
		150.43	Acute on chronic combined systolic (congestive) and diastolic (congestive) heart failure
		150.9	Heart failure, unspecified
		J81.0	Acute pulmonary edema
Diabetes short-term complication	Diabetes short-term complication* (PQI 01)	E1010	Type 1 diabetes mellitus with ketoacidosis without coma
		E1011	Type 1 diabetes mellitus with ketoacidosis with coma
		E10641	Type 1 diabetes mellitus with hypoglycemia with coma
		E1065	Type 1 diabetes mellitus with hyperglycemia
		E1100	Type 2 diabetes mellitus with hyperosmolarity without nonketotic hyperglycemic-hyperosmolar coma (NKHHC)

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		E1101	Type 2 diabetes mellitus with hyperosmolarity with coma
		E11641	Type 2 diabetes mellitus with hypoglycemia with coma
		E1165	Type 2 diabetes mellitus with hyperglycemia
		E08.10	Diabetes mellitus due to underlying condition with ketoacidosis without coma
		E09.10	Drug or chemical induced diabetes mellitus with ketoacidosis without coma
		E13.10	Other specified diabetes mellitus with ketoacidosis without coma
		E08.65	Diabetes mellitus due to underlying condition with hyperglycemia
		E08.01	Diabetes mellitus due to underlying condition with hyperosmolarity with coma
		E09.01	Drug or chemical induced diabetes mellitus with hyperosmolarity with coma
		E13.00	Other specified diabetes mellitus with hyperosmolarity without nonketotic hyperglycemic-hyperosmolar coma (NKHHC)
		E08.65	Diabetes mellitus due to underlying condition with hyperglycemia
		E08.11	Diabetes mellitus due to underlying condition with ketoacidosis with coma
		E08.641	Diabetes mellitus due to underlying condition with hypoglycemia with coma
		E09.11	Drug or chemical induced diabetes mellitus with ketoacidosis with coma
		E09.641	Drug or chemical induced diabetes mellitus with hypoglycemia with coma
		E13.11	Other specified diabetes mellitus with ketoacidosis with coma
		E13.641	Other specified diabetes mellitus with hypoglycemia with coma
		E09.65	Drug or chemical induced diabetes mellitus with hyperglycemia
		E08.618	Diabetes mellitus due to underlying condition with other diabetic arthropathy
		E08.620	Diabetes mellitus due to underlying condition with diabetic dermatitis

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		E08.621	Diabetes mellitus due to underlying condition with foot ulcer
		E08.622	Diabetes mellitus due to underlying condition with other skin ulcer
		E08.628	Diabetes mellitus due to underlying condition with other skin complications
		E08.630	Diabetes mellitus due to underlying condition with periodontal disease
		E08.638	Diabetes mellitus due to underlying condition with other oral complications
		E08.65	Diabetes mellitus due to underlying condition with hyperglycemia
		E08.69	Diabetes mellitus due to underlying condition with other specified complication
		E09.618	Drug or chemical induced diabetes mellitus with other diabetic arthropathy
		E09.621	Drug or chemical induced diabetes mellitus with foot ulcer
		E09.622	Drug or chemical induced diabetes mellitus with other skin ulcer
		E09.628	Drug or chemical induced diabetes mellitus with other skin complications
		E09.630	Drug or chemical induced diabetes mellitus with periodontal disease
		E09.638	Drug or chemical induced diabetes mellitus with other oral complications
		E09.649	Drug or chemical induced diabetes mellitus with hypoglycemia without coma
		E09.65	Drug or chemical induced diabetes mellitus with hyperglycemia
		E09.69	Drug or chemical induced diabetes mellitus with other specified complication
		E13.620	Other specified diabetes mellitus with diabetic dermatitis
		E13.621	Other specified diabetes mellitus with foot ulcer
		E13.622	Other specified diabetes mellitus with other skin ulcer
		E13.628	Other specified diabetes mellitus with other skin complications
		E13.638	Other specified diabetes mellitus with other oral complications

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		E13.649	Other specified diabetes mellitus with
			hypoglycemia without coma
		E13.65	Other specified diabetes mellitus with
			hyperglycemia
		E13.69	Other specified diabetes mellitus with
			other specified complication
		E09.69	Drug or chemical induced diabetes
			mellitus with other specified
			complication
		E11.618	Type 2 diabetes mellitus with other
			diabetic arthropathy
		E11.620	Type 2 diabetes mellitus with diabetic
			dermatitis
		E11.621	Type 2 diabetes mellitus with foot ulcer
		E11.622	Type 2 diabetes mellitus with other skin
			ulcer
		E11.628	Type 2 diabetes mellitus with other skin
			complications
		E11.630	Type 2 diabetes mellitus with
			periodontal disease
		E11.638	Type 2 diabetes mellitus with other oral
			complications
		E11.649	Type 2 diabetes mellitus with
			hypoglycemia without coma
		E11.65	Type 2 diabetes mellitus with
			hyperglycemia
		E11.69	Type 2 diabetes mellitus with other
			specified complication
		E10.618	Type 1 diabetes mellitus with other
		710.120	diabetic arthropathy
		E10.620	Type 1 diabetes mellitus with diabetic
		710.121	dermatitis
		E10.621	Type 1 diabetes mellitus with foot ulcer
		E10.622	Type 1 diabetes mellitus with other skin
		F10.620	ulcer
		E10.628	Type 1 diabetes mellitus with other skin
		F10 (20	complications
		E10.630	Type 1 diabetes mellitus with
		E10.620	periodontal disease
		E10.638	Type 1 diabetes mellitus with other oral
		E10.640	complications
		E10.649	Type 1 diabetes mellitus with
		F10.65	hypoglycemia without coma
		E10.65	Type 1 diabetes mellitus with
			hyperglycemia

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		E10.69	Type 1 diabetes mellitus with other
			specified complication
Hypotension/ Hypertension	Hypotension	I95.1	Orthostatic hypotension
		I95.89	Other hypotension
		I95.2	Hypotension due to drugs
		I95.81	Postprocedural hypotension
		I95.89	Other hypotension
		I95.9	Hypotension, unspecified
	Hypertension* (PQI 07)	I10	Essential (primary) hypertension
		I119	Hypertensive heart disease without heart failure
		I129	Hypertensive chronic kidney disease with stage 1 through stage 4 chronic kidney disease, or unspecified chronic kidney disease
		11310	Hypertensive heart and chronic kidney disease without heart failure, with stage 1 through stage 4 chronic kidney disease, or unspecified chronic kidney disease
Influenza		J11.00	Influenza due to unidentified influenza virus with unspecified type of pneumonia
		J12.9	Viral pneumonia, unspecified
		J10.1	Influenza due to other identified influenza virus with other respiratory manifestations
		J11.1	Influenza due to unidentified influenza virus with other respiratory manifestations
		J11.2	Influenza due to unidentified influenza virus with gastrointestinal manifestations
		J11.81	Influenza due to unidentified influenza virus with encephalopathy
		J11.89	Influenza due to unidentified influenza virus with other manifestations
		J09.X1	Influenza due to identified novel influenza A virus with pneumonia
		J09.X2	Influenza due to identified novel influenza A virus with other respiratory manifestations
		J09.X3	Influenza due to identified novel influenza A virus with gastrointestinal manifestations

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		J09.X9	Influenza due to identified novel
			influenza A virus with other
			manifestations
		J10.08	Influenza due to other identified
			influenza virus with other specified
			pneumonia
Bacterial	Bacterial pneumonia* (PQI	J13	Pneumonia due to Streptococcus
pneumonia	11)		pneumoniae
		J14	Pneumonia due to Hemophilus
			influenzae
		J15211	Pneumonia due to Methicillin
			susceptible Staphylococcus
			aureus
		J15212	Pneumonia due to Methicillin resistant
			Staphylococcus
			aureus
		J153	Pneumonia due to streptococcus, group
			B
		J154	Pneumonia due to other streptococci
		J157	Pneumonia due to Mycoplasma
			pneumoniae
		J159	Unspecified bacterial pneumonia
		J160	Chlamydial pneumonia
		J168	Pneumonia due to other specified
		0100	infectious organisms
		J180	Bronchopneumonia, unspecified
		3100	organism
		J181	Lobar pneumonia, unspecified organism
		J188	Other pneumonia, unspecified organism
		J189	Pneumonia, unspecified organism
Urinary tract	Urinary tract infection*	N10	Acute tubulo-interstitial nephritis
infection / Kidney infection	Ormary tract infection	NIO	Acute tuodio-intersutiai nepiirius
miccuon		N119	Chronic tubulo-interstitial nephritis,
		NIII	unspecified
		N12	Tubulo-interstitial nephritis, not
		1112	·
		N151	specified as acute or
		N151 N159	Renal and perinephric abscess
		11139	Renal tubulo-interstitial disease, unspecified
		N16	Renal tubulo-interstitial disorders in
		1110	
		N2004	diseases classified elsewhere
		N2884	Pyelitis cystica
		N2885	Pyeloureteritis cystica
		N2886	Ureteritis cystica

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		N3000	Acute cystitis without hematuria
		N3001	Acute cystitis with hematuria
		N3090	Cystitis, unspecified without hematuria
		N3091	Cystitis, unspecified with hematuria
		N390	Urinary tract infection, site not specified
	Kidney infection	N30.10	Interstitial cystitis (chronic) without
	, and the second		hematuria
		N30.11	Interstitial cystitis (chronic) with
			hematuria
		N30.20	Other chronic cystitis without hematuria
		N30.21	Other chronic cystitis with hematuria
		N30.80	Other cystitis without hematuria
		N30.81	Other cystitis with hematuria
		N34.0	Urethral abscess
C. difficile		A04.7	Enterocolitis due to Clostridium difficile
infection [135			
subset]			
Septicemia		A02.1	Salmonella sepsis
(except in labor)			
[2]			
		A20.7	Septicemic plague
		A22.7	Anthrax sepsis
		A39.4	Meningococcemia, unspecified
		A40.9	Streptococcal sepsis, unspecified
		A41.2	Sepsis due to unspecified
		A 41 O1	staphylococcus
		A41.01	Sepsis due to Methicillin susceptible
		A 41 02	Staphylococcus aureus
		A41.02	Sepsis due to Methicillin resistant
		A41.1	Staphylococcus aureus Sepsis due to other specified
		A41.1	staphylococcus
		A40.3	Sepsis due to Streptococcus pneumoniae
		A41.4	Sepsis due to streptococcus pheumomac  Sepsis due to anaerobes
		A41.50	Gram-negative sepsis, unspecified
		A41.3	Sepsis due to Hemophilus influenzae
		A41.51	Sepsis due to Escherichia coli [E. coli]
		A41.52	Sepsis due to Eschericina con [E. con]  Sepsis due to Pseudomonas
		A4153	Sepsis due to Serratia
		A41.59	Other Gram-negative sepsis
		A41 89	Other specified sensis
		A41.89	Other specified sepsis  Sepsis unspecified organism
		A41.9	Sepsis, unspecified organism

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		R65.20	Severe sepsis without septic shock
		R65.21	Severe sepsis with septic shock
Dehydration/ Electrolyte imbalance [55]	Dehydration*	E860	Dehydration
		E861	Hypovolemia
		E869	Volume depletion, unspecified
	Hyperosmolality and/or hypernatremia~	E870	Hyperosmolality and hypernatremia
	Gastroenteritis~	A080	Rotaviral enteritis
		A0811	Acute gastroenteropathy due to Norwalk agent
		A0819	Acute gastroenteropathy due to other small round
		A082	Aidenoviral enteritis
		A0831	Calicivirus enteritis
		A0832	Astrovirus enteritis
		A0839	Other viral enteritis
		A084	Viral intestinal infection, unspecified
		A088	Other specified intestinal infections
		A09	Infectious gastroenteritis and colitis, unspecified
		K5289	Other specified noninfective gastroenteritis and colitis
		K529	Noninfective gastroenteritis and colitis, unspecified
	Acute kidney failure~	N170	Acute kidney failure with tubular necrosis
		N171	Acute kidney failure with acute cortical necrosis
		N172	Acute kidney failure with medullary necrosis
		N178	Other acute kidney failure
		N179	Acute kidney failure, unspecified
		N19	Unspecified kidney failure
		N990	Postprocedural (acute) (chronic) kidney failure
		E87.2	Acidosis
		E87.3	Alkalosis
		E87.4	Mixed disorder of acid-base balance
		E87.70	Fluid overload, unspecified
		E87.79	Other fluid overload
		E87.5	Hyperkalemia
		E87.6	Hypokalemia
		E87.8	Other disorders of electrolyte and fluid balance, not elsewhere classified

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
Skin and		L03.021	Acute lymphangitis of right finger
subcutaneous			
tissue infections			
[197]			
		L03.022	Acute lymphangitis of left finger
		L03.029	Acute lymphangitis of unspecified
			finger
		L03.041	Acute lymphangitis of right toe
		L03.042	Acute lymphangitis of left toe
		L03.049	Acute lymphangitis of unspecified toe
		L03.121	Acute lymphangitis of right axilla
		L03.122	Acute lymphangitis of left axilla
		L03.123	Acute lymphangitis of right upper limb
		L03.124	Acute lymphangitis of left upper limb
		L03.125	Acute lymphangitis of right lower limb
		L03.126	Acute lymphangitis of left lower limb
		L03.129	Acute lymphangitis of unspecified part of limb
		L03.212	Acute lymphangitis of face
		L03.222	Acute lymphangitis of neck
		L03.321	Acute lymphangitis of abdominal wall
		L03.322	Acute lymphangitis of back [any part
			except buttock]
		L03.323	Acute lymphangitis of chest wall
		L03.324	Acute lymphangitis of groin
		L03.325	Acute lymphangitis of perineum
		L03.326	Acute lymphangitis of umbilicus
		L03.327	Acute lymphangitis of buttock
		L03.329	Acute lymphangitis of trunk,
			unspecified
		L03.891	Acute lymphangitis of head [any part,
			except face]
		L03.898	Acute lymphangitis of other sites
		L03.91	Acute lymphangitis, unspecified
		L03.011	Cellulitis of right finger
		L03.012	Cellulitis of left finger
		L03.019	Cellulitis of unspecified finger
		L03.031	Cellulitis of right toe
		L03.032	Cellulitis of left toe
		L03.039	Cellulitis of unspecified toe
		L03.111	Cellulitis of right axilla
		L03.112	Cellulitis of left axilla
		L03.113	Cellulitis of right upper limb
		L03.114	Cellulitis of left upper limb
		L03.115	Cellulitis of right lower limb
		L03.116	Cellulitis of left lower limb
		L03.119	Cellulitis of unspecified part of limb

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		L03.211	Cellulitis of face
		L03.221	Cellulitis of neck
		L03.311	Cellulitis of abdominal wall
		L03.312	Cellulitis of back [any part except
			buttock]
		L03.313	Cellulitis of chest wall
		L03.314	Cellulitis of groin
		L03.315	Cellulitis of perineum
		L03.316	Cellulitis of umbilicus
		L03.317	Cellulitis of buttock
		L03.319	Cellulitis of trunk, unspecified
		L03.811	Cellulitis of head [any part, except face]
		L03.818	Cellulitis of other sites
		L03.90	Cellulitis, unspecified
		K12.2	Cellulitis and abscess of mouth
		L02.01	Cutaneous abscess of face
		L02.11	Cutaneous abscess of neck
		L02.211	Cutaneous abscess of abdominal wall
		L02.212	Cutaneous abscess of back [any part,
			except buttock]
		L02.213	Cutaneous abscess of chest wall
		L02.214	Cutaneous abscess of groin
		L02.215	Cutaneous abscess of perineum
		L02.216	Cutaneous abscess of umbilicus
		L02.219	Cutaneous abscess of trunk, unspecified
		L02.31	Cutaneous abscess of buttock
		L02.411	Cutaneous abscess of right axilla
		L02.412	Cutaneous abscess of left axilla
		L02.413	Cutaneous abscess of right upper limb
		L02.414	Cutaneous abscess of left upper limb
		L02.415	Cutaneous abscess of right lower limb
		L02.416	Cutaneous abscess of left lower limb
		L02.419	Cutaneous abscess of limb, unspecified
		L02.511	Cutaneous abscess of right hand
		L02.512	Cutaneous abscess of left hand
		L02.519	Cutaneous abscess of unspecified hand
		L02.611	Cutaneous abscess of right foot
		L02.612	Cutaneous abscess of left foot
		L02.619	Cutaneous abscess of unspecified foot
		L02.811	Cutaneous abscess of head [any part,
			except face]
		L02.818	Cutaneous abscess of other sites
		L02.91	Cutaneous abscess, unspecified
		L08.89	Other specified local infections of the
			skin and subcutaneous tissue

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		L08.9	Local infection of the skin and
			subcutaneous tissue, unspecified
Aspiration		J69.0	Pneumonitis due to inhalation of food
pneumonitis;			and vomit
food/vomitus			
[129]			
Arrhythmia		I48.91	Unspecified atrial fibrillation
•		I48.92	Unspecified atrial flutter
		I48.0	Paroxysmal atrial fibrillation
		I48.1	Persistent atrial fibrillation
		I48.3	Typical atrial flutter
		I48.4	Atypical atrial flutter
Intestinal		K56.49	Other impaction of intestine
impaction		120 01.19	outer impaction of intestine
		K56.41	Fecal impaction
Pressure ulcers		L89.90	Pressure ulcer of unspecified site,
555644 0 44004 0			unspecified stage
		L89.009	Pressure ulcer of unspecified elbow,
			unspecified stage
		L89.119	Pressure ulcer of right upper back,
			unspecified stage
		L89.129	Pressure ulcer of left upper back,
			unspecified stage
		L89.139	Pressure ulcer of right lower back,
			unspecified stage
		L89.149	Pressure ulcer of left lower back,
			unspecified stage
		L89.159	Pressure ulcer of sacral region,
			unspecified stage
		L89.209	Pressure ulcer of unspecified hip,
			unspecified stage
		L89.309	Pressure ulcer of unspecified buttock,
			unspecified stage
		L89.509	Pressure ulcer of unspecified ankle,
			unspecified stage
		L89.609	Pressure ulcer of unspecified heel,
			unspecified stage
		L89.819	Pressure ulcer of head, unspecified stage
		L89.899	Pressure ulcer of other site, unspecified
			stage
		L89.000	Pressure ulcer of unspecified elbow,
			unstageable
		L89.003	Pressure ulcer of unspecified elbow,
			stage 3
		L89.004	Pressure ulcer of unspecified elbow,
			stage 4
		L89.010	Pressure ulcer of right elbow,
			unstageable

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		L89.013	Pressure ulcer of right elbow, stage 3
		L89.014	Pressure ulcer of right elbow, stage 4
		L89.019	Pressure ulcer of right elbow,
			unspecified stage
		L89.020	Pressure ulcer of left elbow, unstageable
		L89.023	Pressure ulcer of left elbow, stage 3
		L89.024	Pressure ulcer of left elbow, stage 4
		L89.029	Pressure ulcer of left elbow, unspecified stage
		L89.100	Pressure ulcer of unspecified part of back, unstageable
		L89.103	Pressure ulcer of unspecified part of
		L69.103	back, stage 3
		L89.104	Pressure ulcer of unspecified part of back, stage 4
		L89.109	Pressure ulcer of unspecified part of back, unspecified stage
		L89.110	Pressure ulcer of right upper back, unstageable
		L89.113	Pressure ulcer of right upper back, stage 3
		L89.114	Pressure ulcer of right upper back, stage 4
		L89.120	Pressure ulcer of left upper back, unstageable
		L89.123	Pressure ulcer of left upper back, stage 3
		L89.124	Pressure ulcer of left upper back, stage 4
		L89.130	Pressure ulcer of right lower back, unstageable
		L89.133	Pressure ulcer of right lower back, stage 3
		L89.134	Pressure ulcer of right lower back, stage 4
		L89.140	Pressure ulcer of left lower back, unstageable
		L89.143	Pressure ulcer of left lower back, stage 3
		L89.144	Pressure ulcer of left lower back, stage 4
		L89.150	Pressure ulcer of sacral region, unstageable
		L89.153	Pressure ulcer of sacral region, stage 3
		L89.154	Pressure ulcer of sacral region, stage 4

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		L89.200	Pressure ulcer of unspecified hip,
			unstageable
		L89.203	Pressure ulcer of unspecified hip, stage
			3
		L89.204	Pressure ulcer of unspecified hip, stage
			4
		L89.210	Pressure ulcer of right hip, unstageable
		L89.213	Pressure ulcer of right hip, stage 3
		L89.214	Pressure ulcer of right hip, stage 4
		L89.219	Pressure ulcer of right hip, unspecified
			stage
		L89.220	Pressure ulcer of left hip, unstageable
		L89.223	Pressure ulcer of left hip, stage 3
		L89.224	Pressure ulcer of left hip, stage 4
		L89.229	Pressure ulcer of left hip, unspecified
			stage
		L89.300	Pressure ulcer of unspecified buttock,
			unstageable
		L89.303	Pressure ulcer of unspecified buttock,
			stage 3
		L89.304	Pressure ulcer of unspecified buttock,
			stage 4
		L89.309	Pressure ulcer of unspecified buttock,
			unspecified stage
		L89.310	Pressure ulcer of right buttock,
			unstageable
		L89.313	Pressure ulcer of right buttock, stage 3
		L89.314	Pressure ulcer of right buttock, stage 4
		L89.319	Pressure ulcer of right buttock,
		X 00 220	unspecified stage
		L89.320	Pressure ulcer of left buttock,
		T 00 222	unstageable
		L89.323	Pressure ulcer of left buttock, stage 3
		L89.324	Pressure ulcer of left buttock, stage 4
		L89.329	Pressure ulcer of left buttock,
		T 00 40	unspecified stage
		L89.40	Pressure ulcer of contiguous site of
		1.00.42	back, buttock and hip, unspecified stage
		L89.43	Pressure ulcer of contiguous site of
		1.90.44	back, buttock and hip, stage 3
		L89.44	Pressure ulcer of contiguous site of
		L89.45	back, buttock and hip, stage 4
		L09.43	Pressure ulcer of contiguous site of back, buttock and hip, unstageable
		L89.500	Pressure ulcer of unspecified ankle,
		L09.300	unstageable
			unstageable

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		L89.503	Pressure ulcer of unspecified ankle,
			stage 3
		L89.504	Pressure ulcer of unspecified ankle,
			stage 4
		L89.509	Pressure ulcer of unspecified ankle,
			unspecified stage
		L89.510	Pressure ulcer of right ankle,
			unstageable
		L89.513	Pressure ulcer of right ankle, stage 3
		L89.514	Pressure ulcer of right ankle, stage 4
		L89.519	Pressure ulcer of right ankle,
			unspecified stage
		L89.520	Pressure ulcer of left ankle, unstageable
		L89.523	Pressure ulcer of left ankle, stage 3
		L89.524	Pressure ulcer of left ankle, stage 4
		L89.529	Pressure ulcer of left ankle, unspecified
			stage
		L89.600	Pressure ulcer of unspecified heel,
			unstageable
		L89.603	Pressure ulcer of unspecified heel, stage
			3
		L89.604	Pressure ulcer of unspecified heel, stage 4
		L89.610	Pressure ulcer of right heel, unstageable
		L89.613	Pressure ulcer of right heel, stage 3
		L89.614	Pressure ulcer of right heel, stage 4
		L89.619	Pressure ulcer of right heel, unspecified
			stage
		L89.620	Pressure ulcer of left heel, unstageable
		L89.623	Pressure ulcer of left heel, stage 3
		L89.624	Pressure ulcer of left heel, stage 4
		L89.629	Pressure ulcer of left heel, unspecified
			stage
		L89.629	Pressure ulcer of left heel, unspecified
			stage
		L89.810	Pressure ulcer of head, unstageable
		L89.813	Pressure ulcer of head, stage 3
		L89.814	Pressure ulcer of head, stage 4
		L89.890	Pressure ulcer of other site, unstageable
		L89.893	Pressure ulcer of other site, stage 3
		L89.894	Pressure ulcer of other site, stage 4
		L89.90	Pressure ulcer of unspecified site,
			unspecified stage
		L89.93	Pressure ulcer of unspecified site, stage
			3

Table 2-2. Preliminary List of Conditions for Defining Potentially Preventable Hospital Readmissions for 30-Days Post-PAC Discharge with ICD-10 Codes (continued)

Conditions	Subconditions	ICD-10-CM	ICD-10-CM Description
		L89.94	Pressure ulcer of unspecified site, stage 4
		L89.95	Pressure ulcer of unspecified site, unstageable

SOURCE: List of potentially preventable readmission conditions from RTI International with ICD-10-CM (version: April 2016).

NOTES: [###] indicates CCS code; \*AHRQ PQI ICD-10 v5 specifications

To be considered a potentially preventable readmission, diagnosis codes must be the principal diagnosis on the readmission claim, except where noted.

^Principal of acute bronchitis AND secondary of subcondition COPD

~principal ICD-10-CM code with secondary ICD-10-CM code for subcondition dehydration

Readmission Readmission is for bone marrow, Yes kidney, or other organ transplant (Table PR.1) No Readmission is for maintenance Yes chemotherapy or rehabilitation (Table PR.2) No Readmission includes a Yes potentially planned procedure (Table PR.3) No Principal discharge diagnosis of readmission is acute or complication of care (Table PR.4) Yes No **PLANNED UNPLANNED** 

Figure 2-1. Planned Readmission Algorithm Version 3.0 Flowchart

Source: 2015 Version of the HWR Planned Readmission Algorithm

## Planned Readmission Algorithm Version 3.0 Tables – Hospital Wide Readmission Measure

Table 2-3. Procedure Categories that are Always Planned (Version 3.0)\*

<b>Procedure CCS</b>	Description
64	Bone marrow transplant
105	Kidney transplant
134	Cesarean section**
135	Forceps; vacuum; and breech delivery <sup>††</sup>
176	Other organ transplantation

<sup>\*</sup>Corresponds to Table PR. 1, referenced in Figure 2-1

Table 2-4. Diagnosis Categories that are Always Planned (Version 3.0)\*

Diagnosis CCS	Description
45	Maintenance chemotherapy
194	Forceps delivery;;
196	Normal pregnancy and/or delivery§§
254	Rehabilitation

<sup>\*</sup>Corresponds to Table PR. 2, referenced in Figure 2-1

Table 2-5. Potentially Planned Procedure Categories (Version 3.0)\*

<b>Procedure CCS</b>	Description
3	Laminectomy; excision intervertebral disc
5	Insertion of catheter or spinal stimulator and injection into spinal
9	Other OR therapeutic nervous system procedures
10	Thyroidectomy; partial or complete
12	Other therapeutic endocrine procedures
33	Other OR therapeutic procedures on nose; mouth and pharynx
36	Lobectomy or pneumonectomy
38	Other diagnostic procedures on lung and bronchus
40	Other diagnostic procedures of respiratory tract and mediastinum
43	Heart valve procedures
44	Coronary artery bypass graft (CABG)

 Table 2-5. Potentially Planned Procedure Categories (Version 3.0) (continued)

Procedure CCS	Description
45	Percutaneous transluminal coronary angioplasty (PTCA)
47	Diagnostic cardiac catheterization; coronary arteriography
48	Insertion; revision; replacement; removal of cardiac pacemaker or cardioverter/defibrillator
49	Other OR heart procedures
51	Endarterectomy; vessel of head and neck
52	Aortic resection; replacement or anastomosis
53	Varicose vein stripping; lower limb
55	Peripheral vascular bypass
56	Other vascular bypass and shunt; not heart
59	Other OR procedures on vessels of head and neck
62	Other diagnostic cardiovascular procedures
66	Procedures on spleen
67	Other therapeutic procedures; hemic and lymphatic system
74	Gastrectomy; partial and total
78	Colorectal resection
79	Local excision of large intestine lesion (not endoscopic)
84	Cholecystectomy and common duct exploration
85	Inguinal and femoral hernia repair
86	Other hernia repair
99	Other OR gastrointestinal therapeutic procedures
104	Nephrectomy; partial or complete
106	Genitourinary incontinence procedures
107	Extracorporeal lithotripsy; urinary
109	Procedures on the urethra
112	Other OR therapeutic procedures of urinary tract
113	Transurethral resection of prostate (TURP)
114	Open prostatectomy
119	Oophorectomy; unilateral and bilateral
120	Other operations on ovary
124	Hysterectomy; abdominal and vaginal

**Table 2-5. Potentially Planned Procedure Categories (Version 3.0) (continued)** 

Procedure CCS	Description
129	Repair of cystocele and rectocele; obliteration of vaginal vault
132	Other OR therapeutic procedures; female organs
142	Partial excision bone
152	Arthroplasty knee
153	Hip replacement; total and partial
154	Arthroplasty other than hip or knee
157	Amputation of lower extremity
158	Spinal fusion
159	Other diagnostic procedures on musculoskeletal system
166	Lumpectomy; quadrantectomy of breast
167	Mastectomy
169	Debridement of wound; infection or burn
170	Excision of skin lesion
172	Skin graft
ICD-9 Codes	Description
30.1, 30.29, 30.3, 30.4, 31.74, 34.6	Laryngectomy, revision of tracheostomy, scarification of pleura (from Proc CCS 42- Other OR Rx procedures on respiratory system and mediastinum)
38.18	Endarterectomy leg vessel (from Proc CCS 60- Embolectomy and endarterectomy of lower limbs)
55.03, 55.04	Percutaneous nephrostomy with and without fragmentation (from Proc CCS 103-Nephrotomy and nephrostomy)
94.26, 94.27	Electroshock therapy (from Proc CCS 218- Psychological and psychiatric evaluation and therapy)

<sup>\*</sup>Corresponds to Table PR. 3, referenced in Figure 2-1

**Table 2-6. Acute Diagnosis Categories (Version 3.0)\*** 

Diagnosis CCS	Description
1	Tuberculosis
2	Septicemia (except in labor)
3	Bacterial infection; unspecified site
4	Mycoses
5	HIV infection
7	Viral infection
8	Other infections; including parasitic
9	Sexually transmitted infections (not HIV or hepatitis)
54	Gout and other crystal arthropathies
55	Fluid and electrolyte disorders
60	Acute posthemorrhagic anemia
61	Sickle cell anemia
63	Diseases of white blood cells
76	Meningitis (except that caused by tuberculosis or sexually transmitted disease)
77	Encephalitis (except that caused by tuberculosis or sexually transmitted disease)
78	Other CNS infection and poliomyelitis
82	Paralysis
83	Epilepsy; convulsions
84	Headache; including migraine
85	Coma; stupor; and brain damage
87	Retinal detachments; defects; vascular occlusion; and retinopathy
89	Blindness and vision defects
90	Inflammation; infection of eye (except that caused by tuberculosis or sexually transmitted disease)
91	Other eye disorders
92	Otitis media and related conditions
93	Conditions associated with dizziness or vertigo
99	Hypertension with complications
100	Acute myocardial infarction (with the exception of ICD-9 codes 410.x2)

Table 2-6. Acute Diagnosis Categories (Version 3.0) (continued)

Diagnosis CCS	Description
102	Nonspecific chest pain
104	Other and ill-defined heart disease
107	Cardiac arrest and ventricular fibrillation
109	Acute cerebrovascular disease
112	Transient cerebral ischemia
116	Aortic and peripheral arterial embolism or thrombosis
118	Phlebitis; thrombophlebitis and thromboembolism
120	Hemorrhoids
122	Pneumonia (except that caused by TB or sexually transmitted disease)
123	Influenza
124	Acute and chronic tonsillitis
125	Acute bronchitis
126	Other upper respiratory infections
127	Chronic obstructive pulmonary disease and bronchiectasis
128	Asthma
129	Aspiration pneumonitis; food/vomitus
130	Pleurisy; pneumothorax; pulmonary collapse
131	Respiratory failure; insufficiency; arrest (adult)
135	Intestinal infection
137	Diseases of mouth; excluding dental
139	Gastroduodenal ulcer (except hemorrhage)
140	Gastritis and duodenitis
142	Appendicitis and other appendiceal conditions
145	Intestinal obstruction without hernia
146	Diverticulosis and diverticulitis
148	Peritonitis and intestinal abscess
153	Gastrointestinal hemorrhage
154	Noninfectious gastroenteritis
157	Acute and unspecified renal failure
159	Urinary tract infections
165	Inflammatory conditions of male genital organs

Table 2-6. Acute Diagnosis Categories (Version 3.0) (continued)

Diagnosis CCS	Description
168	Inflammatory diseases of female pelvic organs
172	Ovarian cyst
197	Skin and subcutaneous tissue infections
198	Other inflammatory condition of skin
225	Joint disorders and dislocations; trauma-related
226	Fracture of neck of femur (hip)
227	Spinal cord injury
228	Skull and face fractures
229	Fracture of upper limb
230	Fracture of lower limb
232	Sprains and strains
233	Intracranial injury
234	Crushing injury or internal injury
235	Open wounds of head; neck; and trunk
237	Complication of device; implant or graft
238	Complications of surgical procedures or medical care
239	Superficial injury; contusion
240	Burns
241	Poisoning by psychotropic agents
242	Poisoning by other medications and drugs
243	Poisoning by nonmedicinal substances
244	Other injuries and conditions due to external causes
245	Syncope
246	Fever of unknown origin
247	Lymphadenitis
249	Shock
250	Nausea and vomiting
251	Abdominal pain
252	Malaise and fatigue
253	Allergic reactions
259	Residual codes; unclassified

Table 2-6. Acute Diagnosis Categories (Version 3.0) (continued)

Diagnosis CCS	Description
650	Adjustment disorders
651	Anxiety disorders
652	Attention-deficit, conduct, and disruptive behavior disorders
653	Delirium, dementia, and amnestic and other cognitive disorders
656	Impulse control disorders, NEC
658	Personality disorders
660	Alcohol-related disorders
661	Substance-related disorders
662	Suicide and intentional self-inflicted injury
663	Screening and history of mental health and substance abuse codes
670	Miscellaneous disorders
ICD-9 Codes	Description
Acute ICD-9 codes wi	ithin Dx CCS 97: Peri-; endo-; and myocarditis; cardiomyopathy
03282	Diphtheritic myocarditis
03640	Meningococcal carditis nos
03641	Meningococcal pericarditis
03642	Meningococcal endocarditis
03643	Meningococcal myocarditis
07420	Coxsackie carditis nos
07421	Coxsackie pericarditis
07422	Coxsackie endocarditis
07423	Coxsackie myocarditis
11281	Candidal endocarditis
11503	Histoplasma capsulatum pericarditis
11504	Histoplasma capssulatum endocarditis
11513	Histoplasma duboisii pericarditis
11514	Histoplasma duboisii endocarditis
11593	Histoplasmosis pericarditis
11594	Histoplasmosis endocarditis
1303	Toxoplasma myocarditis
3910	Acute rheumatic pericarditis

Table 2-6. Acute Diagnosis Categories (Version 3.0) (continued)

Diagnosis CCS	Description
3911	Acute rheumatic endocarditis
3912	Acute rheumatic myocarditis
3918	Acute rheumatic heart disease nec
3919	Acute rheumatic heart disease nos
3920	Rheumatic chorea w heart involvement
3980	Rheumatic myocarditis
39890	Rheumatic heart disease nos
39899	Rheumatic heart disease nec
4200	Acute pericarditis in other disease
42090	Acute pericarditis nos
42091	Acute idiopath pericarditis
42099	Acute pericarditis nec
4210	Acute/subacute bacterial endocarditis
4211	Acute endocarditis in other diseases
4219	Acute/subacute endocarditis nos
4220	Acute myocarditis in other diseases
42290	Acute myocarditis nos
42291	Idiopathic myocarditis
42292	Septic myocarditis
42293	Toxic myocarditis
42299	Acute myocarditis nec
4230	Hemopericardium
4231	Adhesive pericarditis
4232	Constrictive pericarditis
4233	Cardiac tamponade
4290	Myocarditis nos
Acute ICD-9 codes w	ithin Dx CCS 105: Conduction disorders
4260	Atrioventricular
42610	Atrioventricular block nos
42611	Atrioventricular block-1st degree
42612	Atrioventricular block-mobitz ii

Table 2-6. Acute Diagnosis Categories (Version 3.0) (continued)

Diagnosis CCS	Description
42613	Atrioventricular block-2nd degree nec
4262	Left bundle branch hemiblock
4263	Left bundle branch block nec
4264	Right bundle branch block
42650	Bundle branch block nos
42651	Right bundle branch block/left posterior fascicular block
42652	Right bundle branch block/left ant fascicular block
42653	Bilateral bundle branch block nec
42654	Trifascicular block
4266	Other heart block
4267	Anomalous atrioventricular excitation
42681	Lown-ganong-levine syndrome
42682	Long qt syn
4269	Conduction
Acute ICD-9 codes wi	ithin Dx CCS 106: Dysrhythmia
4272	Paroxysmal tachycardia nos
7850	Tachycardia nos
42789	Cardiac dysrhythmias nec
4279	Cardiac dysrhythmia noc
42769	Premature beats nec
Acute ICD-9 codes wi	ithin Dx CCS 108: Congestive heart failure; nonhypertensive
39891	Rheumatic heart failure
4280	Congestive heart failure
4281	Left heart failure
42820	Unspecified systolic heart failure
42821	Acute systolic heart failure
42823	Acute on chronic systolic heart failure
42830	Unspecified diastolic heart failure
42831	Acute diastolic heart failure
42833	Acute on chronic diastolic heart failure
42840	Unspec combined syst & dias heart failure

Table 2-6. Acute Diagnosis Categories (Version 3.0) (continued)

Diagnosis CCS	Description
42841	Acute combined systolic & diastolic heart failure
42843	Acute on chronic combined systolic & diastolic heart failure
4289	Heart failure nos
Acute ICD-9 codes w	ithin Dx CCS 149: Biliary tract disease
5740	Calculus of gallbladder with acute cholecystitis
57400	Calculus of gallbladder with acute cholecystitis without mention of obstruction
57401	Calculus of gallbladder with acute cholecystitis with obstruction
5743	Calculus of bile duct with acute cholecystitis
57430	Calculus of bile duct with acute cholecystitis without mention of obstruction
57431	Calculus of bile duct with acute cholecystitis with obstruction
5746	Calculus of gallbladder and bile duct with acute cholecystitis
57460	Calculus of gallbladder with acute cholecystitis without mention of obstruction
57461	Calculus of gallbladder and bile duct with acute cholecystitis with obstruction
5748	Calculus of gallbladder and bile duct with acute and chronic cholecystitis
57480	Calculus of gallbladder obstruction and bile duct with acute and chronic cholecystitis without mention of obstruction
57481	Calculus of gallbladder and bile duct with acute and chronic cholecystitis with obstruction
5750	Acute cholecystitis
57512	Acute and chronic cholecystitis
5761	Cholangitis
Acute ICD-9 codes w	ith Dx CCS 152: Pancreatic disorders
5770	Acute pancreatitis

<sup>\*</sup>Corresponds to Table PR. 4, referenced in Figure 2-1

Source: 2015 Version of the HWR Planned Readmission Algorithm

Table 2-7. AHRQ CCS Single Level Procedure Codes and ICD-9 Procedure Codes Added to Yale's Planned Readmission Algorithm, for the Post-Acute Care Setting

AHRQ CCS Single Level Procedures Codes	Description	Comment
37	Diagnostic Bronchoscopy and Biopsy of Bronchus	
71	Gastrostomy: temporary and permanent	
82	Endoscopic retrograde cannulation of pancreases (ERCP)	
87	Laparoscopy (GI only)	
89	Exploratory Laparotomy	
160	Other therapeutic procedure on muscles and tendons	
164	Other OR therapeutic procedures on musculoskeletal system	
171	Suture of skin and subcutaneous tissue ICD-9	

ICD-9 Procedure Codes	Description	Comment					
Topic: Amputation of	Topic: Amputation of Lower Extremity						
83.82	Graft of muscle or fascia						
86.87	Fat graft of skin and subcutaneous tissue	Required, Diagnosis V58.41, encounter for planned postoperative wound closure					
Topic: Amputation of	of Upper Extremity						
84.1	Upper limb amputation, not otherwise specified						
84.2	Amputation and disarticulation of finger						
84.3	Amputation and disarticulation of thumb						
84.4	Amputation through hand						
84.5	Disarticulation of wrist						
84.6	Amputation through forearm						
84.7	Disarticulation of elbow						

Table 2-7. AHRQ CCS Single Level Procedure Codes and ICD-9 Procedure Codes Added to Yale's Planned Readmission Algorithm, for the Post-Acute Care Setting (continued)

ICD-9 Procedure Codes	Description	Comment
84.8	Amputation through humerus	
84.9	Disarticulation of shoulder	
84.10	Interthoracoscapular amputation	
Topic: Removal of	Vascular Obstruction, Non-Coronary	
38.18	Endarterectomy, lower limb vessels	
38.08	Embolectomy, lower limb arteries	
39.50	Angioplasty or atherectomy of other non- coronary vessels	
00.55	Insertion of drug-eluting stent(s) of other peripheral vessel(s)	
00.60	Insertion of drug-eluting stent(s) of superficial femoral artery	
39.90	Insertion of non-drug-eluting peripheral (non- coronary) vessel stent(s)	
Topic: Colon and Re	ectal Procedures, Selected	
46.85	Dilation of intestine (includes endosopic approach)	
96.8	Insertion of naso-intestinal tube (includes for decompression)	
96.9	Insertion of rectal tube	
46.50	Closure of intestinal stoma, not otherwise specified	Required, Diagnosis code V55.2, attention to ileostomy, and V55.3, attention to colostomy
46.51	Closure of stoma of small intestine	Required, Diagnosis code V55.2, attention to ileostomy, and V55.3, attention to colostomy
46.52	Closure of stoma of large intestine	Required, Diagnosis code V55.2, attention to ileostomy, and V55.3, attention to colostomy

Table 2-7. AHRQ CCS Single Level Procedure Codes and ICD-9 Procedure Codes Added to Yale's Planned Readmission Algorithm, for the Post-Acute Care Setting (continued)

ICD-9 Procedure Codes	Description	Comment
46.86	Endoscopic insertion of colonic stent(s)	
46.87	Other insertion of colonic stent (s)	
Topic: Insertion of I	Feeding Tubes	
44.39	Other gastroenterostomy (GJ-tube)	
46.39	Other enterostomy (J-tube)	
Topic: Routine Devi	ce Replacement	
86.06	Insertion of totally implanted infusion pump	
Topic: Routine Rem	oval of Devices	
84.57	Removal of (cement) spacer (includes antibiotic impregnated spacer)	
97.41	Removal of thoracotomy tube or pleural cavity drain (non-incisional)	
02.43	Removal of ventricular shunt	
97.37	Removal of tracheostomy tube (non-incisional)	
01.27	removal of catheter(s) from cranial cavity or tissue	
86.05	Incision with removal of foreign body or device from skin and subcutaneous tissue	
02.95	Removal of skull tongs or halo traction device	
78.60-78.69	Removal of implanted devices from bone (includes internal and external fixation	
80.00-80.09	Orthopedic implants arthrotomy for removal of prosthesis without replacement	
Topic: Pleuroscleros	sis_	
34.6	Scarification of pleura	
34.92	Injection into thoracic cavity	
Topic: Colon and Re	ectal Procedures, Selected	
51.14	Other close (endoscopic) biopsy of biliary duct or sphincter of Oddi	
51.64	Endoscopic excision or destruction of lesion of biliary ducts or sphincter of Oddi	

Table 2-7. AHRQ CCS Single Level Procedure Codes and ICD-9 Procedure Codes Added to Yale's Planned Readmission Algorithm, for the Post-Acute Care Setting (continued)

ICD-9 Procedure Codes	Description	Comment
51.84	Endoscopic dilation of ampulla and biliary duct	This code became available in CY 2010
51.84	Endoscopic sphincterotomy and papillotomy	
51.85	Endoscopic insertion of nasobiliary drainage tube	
51.86	Endoscopic insertion of stent (tube) into bile duct	
51.87	Endoscopic removal of stone(s)from biliary tract	
Topic: Fistula		
42.84	Repair of esophageal fistula, not elsewhere classified	
44.63	Closure of other gastric fistula (include gastrocolic, gastrojejunocolic fistula)	
46.72	Closure of fistula of duodenum	
46.74	Closure of fistula of small intestine, except duodenum (includes enterocutaneous)	
46.76	Closure of fistula of large intestine	
47.92	Closure of appendiceal fistula	
48.73	Closure of other rectal fistula	
48.93	Repair of perirectal fistula	
49.11	Anal fistulotomy	
49.12	Anal fistulectomy	
49.73	Closure of anal fistula	
19.9	Other repair of middle ear (includes closure of mastoid fistula	
20.93	Repair of oval and round windows (includes closure of fistula)	
21.82	Closure of nasal fistula	
31.62	Closure of fistula of larynx (includes laryngotracheal)	
31.73	Closure of other fistula of trachea (includes tracheoesophageal)	

Table 2-7. AHRQ CCS Single Level Procedure Codes and ICD-9 Procedure Codes Added to Yale's Planned Readmission Algorithm, for the Post-Acute Care Setting (continued)

ICD-9 Procedure Codes	Description	Comment
33.42	Closure of bronchial fistula (includes bronchocutaneous, bronchoesophageal, bronchovisceral)	
34.73	Closure of other fistula of thorax (includes bronchopleural, bronchopleurocutaneous, bronchopleuromediastinal)	
34.83	Closure of fistula of diaphragm (includes thoracicoabdominal, thoracicogastric, thoracicointestinal)	
34.93	Repair of pleura (includes closure of unspecified pleural fistula)	
61.42	Repair of scrotal fistula	
Topic: Tendon Repa	ir (eye)	
15.7	Repair of injury of extraocular muscle (includes repair of tendon)	
Topic: Aneurysm		
39.51	Clipping of aneurysm	

NOTE: December, 2012 Yale added several additional AHRQ CCS Single-Level Procedure Codes. Two of these codes 169 (Debridement of wound; infection or burn) and 172 (Skin graft) had been on the prior RTI developed list.

## **Preliminary Testing Results for the SNF Setting**

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
Variable Name in Woder	Age-Sex Groups (Reference group: Male 18-64)	Count	Total	Estillate	Ellol	1 value	Katios	CL	Opper CL
m65_69	Male age 65-69	55,599	4.0	0.069	0.024	0.005	1.072	1.021	1.126
m70_74	Male age 70-74	62,814	4.5	0.123	0.025	<.0001	1.131	1.076	1.189
m75_79	Male age 75-79	76,329	5.5	0.138	0.024	<.0001	1.149	1.094	1.207
m80_84	Male age 80-84	89,529	6.5	0.210	0.024	<.0001	1.232	1.174	1.293
m85_89	Male age 85-89	87,594	6.3	0.230	0.024	<.0001	1.257	1.196	1.320
m90_94	Male age 90-94	49,799	3.6	0.234	0.027	<.0001	1.262	1.195	1.333
m95_GT	Male age 95+	13,899	1.0	0.278	0.039	<.0001	1.315	1.218	1.421
w18_64	Female age 18-64	58,174	4.2	0.049	0.024	0.039	1.052	1.003	1.103
w65_69	Female age 65-69	76,859	5.6	0.080	0.023	0.001	1.085	1.036	1.137
w70_74	Female age 70-74	100,049	7.2	0.105	0.024	<.0001	1.113	1.062	1.167
w75_79	Female age 75-79	130,692	9.5	0.131	0.023	<.0001	1.141	1.089	1.195
w80_84	Female age 80-84	165,967	12.0	0.146	0.023	<.0001	1.157	1.105	1.212
w85_89	Female age 85-89	184,663	13.4	0.156	0.023	<.0001	1.168	1.116	1.224
w90_94	Female age 90-94	126,126	9.1	0.165	0.024	<.0001	1.177	1.121	1.235
w95_GT	Female age 95+	47,463	3.4	0.129	0.029	<.0001	1.135	1.071	1.202

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
	CCS GROUPS Reference group is Miscellaneous Negative: Immunization (10); Sickle Cell Anemia (61); Cataract (86); Inflammation; infection of Eye (exc. Caused by Tuberculosis or STD) (90); Other Complications of Pregnancy (181); Other Complications of Birth, Puerperium Affecting Management of Mother (195); Lymphadenitis (247); Shock (249); Rehabilitation care; fitting of prostheses; and adjustment of device (254); Medical Exam/Eval (256); Personality Disorders (658)								
Pr_Cond_sngl_2	Septicemia (except in labor) (2)	93,697	6.8	0.971	0.039	<.0001	2.650	2.449	2.868
Pr_Cond_sngl_4	Mycoses (4)	1,508	0.1	1.030	0.091	<.0001	2.814	2.345	3.376
Pr_Cond_sngl_6	Hepatitis (6)	750	0.1	0.657	0.148	<.0001	1.934	1.439	2.600
Viral_Tonsil_Cond	Viral Infection and Tonsillitis: Viral Infection (7); Acute and Chronic Tonsillitis (124)	1,442	0.1	0.877	0.110	<.0001	2.421	1.940	3.022
Bact_Infection_STI_C	Miscellaneous Infections: Bacterial Infection (3), Unspecified Site; Other Infections, Including Parasitic (8); Sexually transmitted infections (not HIV or hepatitis) (9)	774	0.1	0.817	0.152	<.0001	2.302	1.697	3.123
Pr_Cond_sngl_11	Cancer of Head and Neck (11)	563	0.0	1.253	0.202	<.0001	3.516	2.346	5.270
Pr_Cond_sngl_14	Cancer of Colon (14)	4,795	0.3	0.669	0.083	<.0001	1.954	1.655	2.308
Pr_Cond_sngl_15	Cancer of Rectum and Anus (15)	1,186	0.1	1.001	0.136	<.0001	2.715	2.065	3.570
Pr_Cond_sngl_17	Cancer of Pancreas (17)	458	0.0	0.926	0.210	<.0001	2.507	1.646	3.819
Pr_Cond_sngl_19	Cancer of Bronchus, Lung (19)	1,715	0.1	1.182	0.110	<.0001	3.281	2.630	4.094
Pr_Cond_sngl_33	Cancer of Kidney and Renal Pelvis (33)	815	0.1	0.652	0.165	<.0001	1.911	1.372	2.662
Pr_Cond_sngl_35	Cancer of Brain and Nervous System (35)	445	0.0	0.789	0.251	0.002	2.196	1.327	3.632
Pr_Cond_sngl_42	Secondary Malignancies (42)	1,772	0.1	0.878	0.117	<.0001	2.408	1.904	3.044
Gen_Cancer_Cond	General Cancer Conditions: Esophagus (12); Stomach (13); Other GI Organs (18), Peritoneum; Breast (24)	1,564	0.1	0.650	0.136	<.0001	1.908	1.451	2.509

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
Reprod_Cancer_Cond	Female Reproductive Cancers: Uterus (25); Ovary (27); Other Female Genital Organs (28)	1,240	0.1	1.172	0.177	<.0001	3.229	2.261	4.610
Urinary_Cancer_Cond	Urinary System and Prostate Cancers: Prostate (29); Bladder (32); Other Urinary Organs (34)	1,575	0.1	1.167	0.097	<.0001	3.229	2.657	3.924
Unspec_Cancer_Cond	Unspecified Cancer Conditions: Liver and Intrahepatic Bile Duct (16); Hodgkin's Disease (37); Non-Hodgkin's Lymphoma (38); Leukemia (39); Multiple Myeloma (40); Cancer, Other and Unspecified Primary (40); Malignant Neoplasm Without Specification of Site (43); Neoplasms of Unspecified Nature or Uncertain Behavior (44)	1,084	0.1	1.221	0.123	<.0001	3.388	2.645	4.339
Misc_Cancer_Cond	Miscellaneous Cancer Conditions: Other Respiratory and Intrathoracic (20); Bone and Connective Tissue (21); Melanomas of Skin (22); Other Non-Epithelial Cancer of Skin (23); Cervix (26); Testis (30); Male Genital Organs (31); Thyroid (36); Maintenance Chemotherapy, Radiotherapy (45)	782	0.1	0.698	0.190	0.000	2.005	1.368	2.940
Benign_Unspec_Neopla	Benign Neoplasms: Neoplasm of Uterus (46); Other Benign Neoplasm (47)	2,419	0.2	0.655	0.109	<.0001	1.925	1.547	2.396
Pr_Cond_sngl_48	Thyroid disorders (48)	784	0.1	0.954	0.144	<.0001	2.614	1.957	3.491
Diabetes_Cond	Diabetes: Diabetes Without Complication (49); Diabetes With Complication (50)	18,659	1.3	1.010	0.043	<.0001	2.752	2.527	2.998
Pr_Cond_sngl_51	Other endocrine disorders (51)	3,857	0.3	0.800	0.074	<.0001	2.223	1.914	2.581
Pr_Cond_sngl_52	Nutritional deficiencies (52)	750	0.1	0.706	0.158	<.0001	2.032	1.479	2.793
Immune_Metabolic_Dis	Disorders of Lipid Metabolism (53); Immunity Disorders (57); Other Nutritional, Endocrine, and Metabolic Disorders (58)	3,608	0.3	0.897	0.073	<.0001	2.457	2.120	2.846
Pr_Cond_sngl_54	Gout and other crystal arthropathies (54)	1,200	0.1	0.767	0.124	<.0001	2.160	1.684	2.770
Pr_Cond_sngl_55	Fluid and electrolyte disorders (55)	22,840	1.7	1.016	0.042	<.0001	2.779	2.556	3.021
CysticFibr_Cond	Cystic Fibrosis, COPD and Bronchiectasis: Cystic Fibrosis (56); COPD and Bronchiectasis (127)	30,317	2.2	1.416	0.038	<.0001	4.145	3.842	4.472

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
Pr_Cond_sngl_59	Deficiency and Other Anemia (59)	8,388	0.6	0.881	0.052	<.0001	2.420	2.179	2.688
Pr_Cond_sngl_60	Acute post hemorrhagic anemia (60)	2,622	0.2	0.730	0.084	<.0001	2.076	1.755	2.456
Coag_Hemotal_Cond	Coagulation, Hemorrhagic and Other Blood Disorders: Coagulation and Hemorrhagic Disorder (62); Other Hematologic Conditions (64)	865	0.1	0.755	0.137	<.0001	2.146	1.630	2.825
Pr_Cond_sngl_63	Diseases of White Blood Cells (63)	983	0.1	0.644	0.136	<.0001	1.908	1.453	2.505
Anx_Dep_Cond	Anxiety Disorder (651); Mood Disorder (657); Suicide and Intentional Self-Inflicted Injury (662)	8,926	0.6	0.361	0.064	<.0001	1.417	1.246	1.612
Impulse_Cond	Adjustment Disord (650); Attention Deficit, Conduct and Disruptive Behavior (652); Impulse Ctrl Disord (656)	417	0.0	0.378	0.253	0.135	1.457	0.877	2.419
Pr_Cond_sngl_653	Delirium (653)	23,851	1.7	0.442	0.048	<.0001	1.541	1.400	1.696
Pr_Cond_sngl_659	Schizophrenia and other psychotic disorders (659)	8,570	0.6	0.155	0.072	0.032	1.149	0.994	1.327
Pr_Cond_sngl_660	Alcohol-related disorders (660)	3,766	0.3	0.276	0.094	0.004	1.312	1.086	1.586
Pr_Cond_sngl_661	Substance-related disorders (661)	2,655	0.2	0.658	0.088	<.0001	1.918	1.608	2.289
Pr_Cond_sngl_76	Meningitis (except that caused by tuberculosis or sexually transmitted disease) (76)	451	0.0	0.963	0.189	<.0001	2.598	1.778	3.797
CNS_Infection_Cond	Encephalitis and CNS Infections: Encephalitis (77); Oth CNS infx polio (78)	968	0.1	0.396	0.168	0.018	1.483	1.059	2.077
Pr_Cond_sngl_79	Parkinson's disease (79)	1,649	0.1	0.588	0.130	<.0001	1.809	1.392	2.350
Pr_Cond_sngl_80	Multiple sclerosis (80)	831	0.1	0.861	0.159	<.0001	2.391	1.737	3.292
Pr_Cond_sngl_81	Other hereditary and degenerative nervous system conditions (81)	3,344	0.2	0.660	0.090	<.0001	1.940	1.619	2.325
Pr_Cond_sngl_83	Epilepsy, convulsions (83)	8,517	0.6	0.517	0.061	<.0001	1.676	1.482	1.894

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
Paral_Cereb_LateEff	Paralysis and Late Effects of Cerebrovascular Disease: Paralysis (82); Cereb Late Eff (113)	1,945	0.1	0.544	0.112	<.0001	1.726	1.379	2.160
Coma_Headache_Cond	Headache and Coma: Headache (84); Coma (85)	1,567	0.1	0.572	0.121	<.0001	1.764	1.384	2.249
Eye_Ear_Cond	Sensory Organ Conditions: Retinal (87); Glaucoma (88); Blindness (89); Oth Eye (91); Otitis (92); Oth Ear Sensory Organ (94)	602	0.0	0.904	0.171	<.0001	2.478	1.759	3.491
Pr_Cond_sngl_93	Conditions associated with dizziness or vertigo (93)	2,232	0.2	0.328	0.124	0.008	1.404	1.094	1.802
Pr_Cond_sngl_95	Other nervous system disorders (95)	17,083	1.2	0.719	0.047	<.0001	2.058	1.874	2.260
Pr_Cond_sngl_96	Heart valve disorders (96)	9,866	0.7	0.992	0.059	<.0001	2.695	2.393	3.037
Pr_Cond_sngl_97	Peri- endo- & myocarditis cardiomyopathy (except caused by tuberculosis or sexually transmitted disease) (97)	1,563	0.1	0.562	0.111	<.0001	1.755	1.406	2.191
Pr_Cond_sngl_98	Essential hypertension (98)	1,205	0.1	0.656	0.142	<.0001	1.950	1.467	2.592
Pr_Cond_sngl_99	Hypertension with complications and secondary hypertension (99)	9,091	0.7	1.032	0.048	<.0001	2.827	2.567	3.112
Pr_Cond_sngl_100	Acute myocardial infarction (100)	22,006	1.6	0.971	0.042	<.0001	2.644	2.430	2.877
Atherosclerosis_Cond	Atherosclerosis and Other Heart Disease: Coronary Atherosclerosis and Oth Heart Disease (101); Other and Ill Defined Heart (104)	10,665	0.8	0.799	0.057	<.0001	2.235	1.995	2.505
Pr_Cond_sngl_102	Nonspecific chest pain (102)	3,700	0.3	0.822	0.071	<.0001	2.285	1.983	2.633
Pr_Cond_sngl_103	Pulmonary heart disease (103)	8,438	0.6	0.689	0.057	<.0001	1.989	1.775	2.228
Pr_Cond_sngl_105	Conduction disorders (105)	3,189	0.2	0.616	0.086	<.0001	1.851	1.559	2.199
Pr_Cond_sngl_106	Cardiac dysrhythmias (106)	27,972	2.0	1.080	0.040	<.0001	2.948	2.723	3.193
Pr_Cond_sngl_107	Cardiac arrest and ventricular fibrillation (107)	352	0.0	0.752	0.197	0.000	2.121	1.428	3.150
Pr_Cond_sngl_108	Congestive heart failure; nonhypertensive (108)	56,842	4.1	1.217	0.036	<.0001	3.389	3.151	3.644
Pr_Cond_sngl_109	Acute cerebrovascular disease (109)	48,070	3.5	0.632	0.040	<.0001	1.881	1.735	2.038
Pr_Cond_sngl_110	Occlusion or stenosis of precerebral arteries (110)	1,763	0.1	0.864	0.112	<.0001	2.377	1.900	2.974

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
Pr_Cond_sngl_111	Other and ill-defined cerebrovascular disease (111)	861	0.1	0.358	0.185	0.053	1.419	0.978	2.058
Pr_Cond_sngl_112	Transient cerebral ischemia (112)	6,971	0.5	0.680	0.064	<.0001	1.982	1.743	2.255
Pr_Cond_sngl_114	Peripheral and visceral atherosclerosis (114)	7,145	0.5	0.756	0.061	<.0001	2.132	1.886	2.411
Pr_Cond_sngl_115	Aortic; peripheral; and visceral artery aneurysms (115)	2,936	0.2	0.497	0.103	<.0001	1.640	1.333	2.017
Pr_Cond_sngl_116	Aortic and peripheral arterial embolism or thrombosis (116)	1,725	0.1	0.659	0.115	<.0001	1.938	1.540	2.440
Pr_Cond_sngl_117	Other circulatory disease (117)	7,334	0.5	0.834	0.057	<.0001	2.312	2.063	2.590
Pr_Cond_sngl_118	Phlebitis; thrombophlebitis and thromboembolism (118)	7,920	0.6	0.708	0.058	<.0001	2.039	1.813	2.293
Varicose_Hemmr_Cond	Other Vein: Varicose (119); Hemorrhoids (120)	1,097	0.1	0.791	0.122	<.0001	2.216	1.735	2.831
Pr_Cond_sngl_121	Other diseases of veins and lymphatics (121)	1,456	0.1	0.905	0.104	<.0001	2.478	2.012	3.052
Pr_Cond_sngl_122	Pneumonia (except that caused by tuberculosis or sexually transmitted disease) (122)	65,999	4.8	1.036	0.036	<.0001	2.825	2.626	3.039
Pr_Cond_sngl_123	Influenza (123)	4,668	0.3	0.673	0.072	<.0001	1.958	1.693	2.264
Pr_Cond_sngl_125	Acute bronchitis (125)	2,910	0.2	0.813	0.086	<.0001	2.269	1.910	2.697
Pr_Cond_sngl_126	Other upper respiratory infections (126)	694	0.1	0.926	0.157	<.0001	2.535	1.849	3.475
Pr_Cond_sngl_128	Asthma (128)	4,782	0.3	1.235	0.057	<.0001	3.472	3.095	3.894
Pr_Cond_sngl_129	Aspiration pneumonitis; food/vomitus (129)	17,642	1.3	1.039	0.043	<.0001	2.827	2.595	3.079
Pr_Cond_sngl_130	Pleurisy; pneumothorax; pulmonary collapse (130)	4,238	0.3	0.857	0.066	<.0001	2.355	2.062	2.688
Pr_Cond_sngl_131	Respiratory failure; insufficiency; arrest (adult) (131)	18,995	1.4	0.955	0.042	<.0001	2.589	2.382	2.815
RespDisease_Cond	Lung Disease Due to External Agents and Other Upper Respiratory Disease: Lung Disease External (132); Other Upper Resp (134)	1,140	0.1	0.907	0.113	<.0001	2.484	1.981	3.114
Pr_Cond_sngl_133	Other lower respiratory disease (133)	2,828	0.2	0.793	0.079	<.0001	2.211	1.887	2.590
Pr_Cond_sngl_135	Intestinal infection (135)	11,993	0.9	1.426	0.043	<.0001	4.187	3.839	4.565
MouthDisease_Cond	Disease and Disorder of Mouth, Teeth, and Jaw: Disorder Teeth Jaw (136); Disease Mouth Excl Dent (137)	774	0.1	0.717	0.154	<.0001	2.041	1.499	2.779

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
Pr_Cond_sngl_138	Esophageal disorders (138)	3,086	0.2	0.666	0.082	<.0001	1.949	1.652	2.301
Pr_Cond_sngl_139	Gastroduodenal ulcer (except hemorrhage) (139)	1,408	0.1	0.816	0.121	<.0001	2.273	1.782	2.899
Gastric_Cond	Gastritis Duodenitis (140); Peritonitis intestinal abscess (148)	3,115	0.2	0.672	0.080	<.0001	1.965	1.672	2.308
Pr_Cond_sngl_141	Other disorders of stomach and duodenum (141)	2,071	0.1	0.644	0.094	<.0001	1.900	1.575	2.292
Pr_Cond_sngl_142	Appendicitis and other appendiceal conditions (142)	1,021	0.1	0.575	0.173	0.001	1.778	1.256	2.516
Pr_Cond_sngl_143	Abdominal hernia (143)	6,072	0.4	0.655	0.074	<.0001	1.924	1.658	2.233
Pr_Cond_sngl_144	Regional enteritis and ulcerative colitis (144)	867	0.1	1.298	0.118	<.0001	3.696	2.917	4.682
Pr_Cond_sngl_145	Intestinal obstruction without hernia (145)	13,264	1.0	0.516	0.053	<.0001	1.677	1.506	1.866
Pr_Cond_sngl_146	Diverticulosis and diverticulitis (146)	8,860	0.6	0.627	0.059	<.0001	1.876	1.667	2.112
Pr_Cond_sngl_147	Anal and rectal conditions (147)	1,590	0.1	0.997	0.106	<.0001	2.733	2.210	3.379
Pr_Cond_sngl_149	Biliary tract disease (149)	9,214	0.7	0.752	0.058	<.0001	2.127	1.892	2.392
Pr_Cond_sngl_151	Other liver diseases (151)	3,766	0.3	0.778	0.071	<.0001	2.177	1.889	2.509
Pr_Cond_sngl_152	Pancreatic disorders (not diabetes) (152)	3,634	0.3	0.600	0.081	<.0001	1.828	1.553	2.152
Pr_Cond_sngl_153	Gastrointestinal hemorrhage (153)	18,207	1.3	0.743	0.045	<.0001	2.100	1.920	2.296
Pr_Cond_sngl_154	Noninfectious gastroenteritis (154)	2,947	0.2	0.951	0.078	<.0001	2.617	2.238	3.061
Pr_Cond_sngl_155	Other gastrointestinal disorders (155)	7,026	0.5	0.722	0.060	<.0001	2.061	1.827	2.324
Nephritis_OtherKidne	Nephritis and Other Kidney: Nephritis, Nephrosis, Renal Sclerosis (156); Other Dis Kidney Ureter (161)	1,123	0.1	1.041	0.110	<.0001	2.826	2.268	3.523
Pr_Cond_sngl_157	Acute and unspecified renal failure (157)	40,556	2.9	1.095	0.038	<.0001	3.002	2.783	3.238
Pr_Cond_sngl_158	Chronic renal failure (158)	471	0.0	0.808	0.159	<.0001	2.241	1.629	3.084
Pr_Cond_sngl_159	Urinary tract infections (159)	56,036	4.1	1.046	0.037	<.0001	2.867	2.661	3.089
Pr_Cond_sngl_160	Calculus of urinary tract (160)	1,367	0.1	0.914	0.107	<.0001	2.496	2.015	3.091
Pr_Cond_sngl_162	Other diseases of bladder and urethra (162)	902	0.1	0.830	0.135	<.0001	2.285	1.742	2.998

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
Pr_Cond_sngl_163	Genitourinary symptoms and ill-defined conditions (163)	1,554	0.1	0.838	0.102	<.0001	2.318	1.890	2.843
Pr_Cond_sngl_164	Hyperplasia of prostate (164)	1,163	0.1	0.775	0.122	<.0001	2.181	1.705	2.788
Male_Genital_Cond	Male Genital Disorders: Infl cond of male genital org (165); Oth Male Genital (166)	835	0.1	0.831	0.136	<.0001	2.309	1.758	3.032
Female_Reprod_Cond	Female Reproductive Disorders: Infl Fem Pelvic (168); Endometriosis (169); Prolapse Female Gen (170); Menstrual Disorder (171); Ov Cyst (172); Menopausal Disorder (173); Fem Infertility (174); Oth Fem Gen (175)	1,061	0.1	0.772	0.147	<.0001	2.182	1.626	2.928
Pr_Cond_sngl_197	Skin and subcutaneous tissue infections (197)	24,519	1.8	1.152	0.040	<.0001	3.191	2.942	3.461
Skin_Cond	Skin Disorders: Other Inflammatory Condition of Skin (198); Other Skin Disorders (200)	543	0.0	0.793	0.178	<.0001	2.212	1.548	3.159
Pr_Cond_sngl_199	Chronic ulcer of skin (199)	3,968	0.3	0.935	0.069	<.0001	2.565	2.234	2.944
Pr_Cond_sngl_201	Infective arthritis and osteomyelitis (except that caused by tuberculosis or sexually transmitted disease) (201)	5,317	0.4	0.797	0.066	<.0001	2.227	1.949	2.544
Pr_Cond_sngl_202	Rheumatoid arthritis and related disease (202)	954	0.1	0.762	0.162	<.0001	2.166	1.564	2.999
Pr_Cond_sngl_204	Other non-traumatic joint disorders (204)	3,987	0.3	0.538	0.092	<.0001	1.715	1.427	2.063
Pr_Cond_sngl_205	Spondylosis; intervertebral disc disorders; other back problems (205)	24,373	1.8	0.551	0.047	<.0001	1.742	1.585	1.914
Pr_Cond_sngl_207	Pathological Fracture (207)	10,747	0.8	0.858	0.054	<.0001	2.367	2.125	2.637
Pr_Cond_sngl_209	Other acquired deformities (209)	3,390	0.2	0.434	0.120	0.000	1.542	1.211	1.964
Conn_Tissue_Cond	Systemic Lupus and Connective Tissue Disorders: Systemic lupus erythematosus and Connective (210); Other Connective (211)	12,218	0.9	0.716	0.053	<.0001	2.051	1.843	2.282
Bone_Disease_Cond	Osteoporosis and Other Bone Disease: Osteoporosis (206); Oth Bone Disease (212)	4,789	0.3	0.491	0.092	<.0001	1.635	1.358	1.969
Congenital_Cond	Congenital Anomalies: Cardiac/Circulatory (213); Digestive (214); Genitourinary (215); Nervous Syst (216); Other (217)	1,083	0.1	0.662	0.176	0.000	1.927	1.352	2.745

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
Pr_Cond_sngl_225	Joint disorders and dislocations; trauma-related (225)	1,551	0.1	0.494	0.159	0.002	1.642	1.192	2.261
Pr_Cond_sngl_226	Fracture of neck of femur (hip) (226)	95,502	6.9	0.744	0.032	<.0001	2.107	1.975	2.249
Pr_Cond_sngl_227	Spinal cord injury (227)	559	0.0	0.673	0.221	0.002	1.971	1.266	3.070
Pr_Cond_sngl_228	Skull and face fractures (228)	1,463	0.1	0.576	0.138	<.0001	1.779	1.349	2.347
Pr_Cond_sngl_229	Fracture of upper limb (229)	14,741	1.1	0.710	0.052	<.0001	2.044	1.841	2.268
Pr_Cond_sngl_230	Fracture of lower limb (230)	25,644	1.9	0.654	0.044	<.0001	1.931	1.767	2.111
Pr_Cond_sngl_231	Other fractures (231)	35,815	2.6	0.633	0.042	<.0001	1.886	1.734	2.051
Pr_Cond_sngl_232	Sprains and strains (232)	2,171	0.2	0.511	0.119	<.0001	1.672	1.316	2.125
Pr_Cond_sngl_233	Intracranial injury (233)	12,985	0.9	0.571	0.055	<.0001	1.769	1.583	1.977
Pr_Cond_sngl_234	Crushing injury or internal injury (234)	2,736	0.2	0.502	0.098	<.0001	1.650	1.354	2.010
Pr_Cond_sngl_235	Open wounds of head; neck; and trunk (235)	871	0.1	0.851	0.149	<.0001	2.356	1.748	3.176
Pr_Cond_sngl_236	Open wounds of extremities (236)	869	0.1	1.039	0.138	<.0001	2.841	2.151	3.751
Pr_Cond_sngl_237	Complication of device; implant or graft (237)	42,237	3.1	0.609	0.038	<.0001	1.838	1.704	1.983
Pr_Cond_sngl_238	Complications of surgical procedures or medical care (238)	17,800	1.3	0.572	0.046	<.0001	1.770	1.613	1.943
Pr_Cond_sngl_239	Superficial injury; contusion (239)	3,960	0.3	0.585	0.082	<.0001	1.807	1.533	2.129
Ext_Inj_Cond	External Injuries: Burns (240); Other inj cond external (244)	3,333	0.2	0.711	0.083	<.0001	2.040	1.727	2.409
Poisoning_Cond	Poisoning: Psychotropic Agents (241); Other Medications and Drugs (242); Nonmedicinal Substances (243)	2,898	0.2	0.676	0.084	<.0001	1.966	1.662	2.325
Pr_Cond_sngl_245	Syncope (245)	8,029	0.6	0.671	0.061	<.0001	1.975	1.748	2.232
Pr_Cond_sngl_246	Fever of unknown origin (246)	1,267	0.1	0.692	0.124	<.0001	1.992	1.553	2.555
Pr_Cond_sngl_248	Gangrene (248)	3,235	0.2	0.793	0.080	<.0001	2.224	1.893	2.613
Pr_Cond_sngl_250	Nausea and vomiting (250)	899	0.1	0.792	0.136	<.0001	2.213	1.684	2.907
Pr_Cond_sngl_251	Abdominal pain (251)	1,495	0.1	0.601	0.116	<.0001	1.830	1.449	2.310
Pr_Cond_sngl_252	Malaise and fatigue (252)	3,540	0.3	0.763	0.080	<.0001	2.143	1.825	2.515

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
Pr_Cond_sngl_253	Allergic reactions (253)	402	0.0	0.813	0.196	<.0001	2.267	1.528	3.363
Pr_Cond_sngl_259	Residual codes; unclassified (259)	6,606	0.5	0.832	0.060	<.0001	2.300	2.040	2.593
Misc_Pos_PPR	Miscellaneous Positive: Tuberculosis (1); HIV (5); Nonmalignant Breast (167); Menopausal Disorders (173); Acquired Foot Deform (208); Admin/social (255); Other Aftercare (257); Oth Screen (258); Developmental Disord (654); Misc Mental Disord (670)	2,230	0.2	0.326	0.115	0.005	1.382	1.096	1.742
	Surgical Groups								
vascular	Vascular Surgery	10,659	0.8	-0.146	0.052	0.005	0.862	0.777	0.956
neuro	Neurosurgery	11,059	0.8	-0.198	0.056	0.000	0.817	0.729	0.914
ct	Cardio Thoracic Surgery	28,120	2.0	-0.246	0.035	<.0001	0.779	0.727	0.835
Ortho	Orthopedics Surgery	318,323	23.0	-0.408	0.022	<.0001	0.663	0.634	0.692
gen	General surgery	66,769	4.8	-0.276	0.023	<.0001	0.758	0.724	0.793
plastic	Plastic Surgery	20,359	1.5	-0.106	0.035	0.003	0.898	0.838	0.964
ent	Otolaryngology Surgery	2,432	0.2	-0.145	0.100	0.147	0.866	0.709	1.058
obgyn	Obstetrics/Gynecology Surgery	3,465	0.3	-0.348	0.114	0.002	0.704	0.560	0.884
	End Stage Renal Disease								
esrd_indicator	End Stage Renal Disease Indicator	43,352	3.1	0.218	0.029	<.0001	1.245	1.175	1.318
	Comorbidities - Hierarchical Condition Categories - HCCs in year prior)	(Note: * If from	n prior prox	kimal stay; o	otherwise,				
HCC_prox_2	HCC2 Septicemia, Sepsis, Systemic Inflammatory Response Syndrome/Shock *	123,131	8.9	0.030	0.020	0.123	1.031	0.991	1.072
HCC365_6	HCC6 Opportunistic Infections	10,409	0.8	0.096	0.031	0.002	1.101	1.035	1.172
HCC365_7	HCC7 Other Infectious Diseases	347,082	25.1	0.035	0.009	<.0001	1.034	1.016	1.052
HCC365_8	HCC8 Metastatic Cancer and Acute Leukemia	29,329	2.1	0.099	0.023	<.0001	1.104	1.054	1.156

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
HCC365_9	HCC9 Lung and Other Severe Cancers	19,933	1.4	0.081	0.025	0.001	1.087	1.033	1.144
HCC365_10	HCC10 Lymphoma and Other Cancers	19,824	1.4	0.167	0.026	<.0001	1.180	1.120	1.244
HCC365_17	HCC17 Diabetes with Acute Complications	9,294	0.7	0.287	0.035	<.0001	1.332	1.243	1.429
HCC365_18	HCC18 Diabetes with Chronic Complications	140,480	10.2	0.128	0.013	<.0001	1.138	1.109	1.167
HCC365_19	HCC19 Diabetes without Complication	336,201	24.3	0.085	0.009	<.0001	1.092	1.073	1.110
HCC365_20	HCC20 Type I Diabetes Mellitus	12,437	0.9	0.087	0.031	0.005	1.092	1.026	1.161
HCC365_21	HCC21 Protein-Calorie Malnutrition	166,114	12.0	0.030	0.010	0.002	1.029	1.009	1.050
HCC_prox_22	HCC22 Morbid Obesity *	84,316	6.1	0.022	0.014	0.126	1.020	0.991	1.050
HCC365_23	HCC23 Other Significant Endocrine and Metabolic Disorders	70,054	5.1	0.029	0.014	0.047	1.026	0.997	1.056
HCC365_24	HCC24 Disorders of Fluid/Electrolyte/Acid-Base Balance	725,945	52.5	0.079	0.008	<.0001	1.081	1.064	1.100
HCC365_26	HCC26 Other Endocrine/Metabolic/Nutritional Disorders	565,553	40.9	0.013	0.007	0.074	1.011	0.996	1.026
HCC_prox_35	HCC35 Inflammatory Bowel Disease *	7,838	0.6	0.080	0.043	0.062	1.078	0.989	1.175
HCC_prox_36	HCC36 Peptic Ulcer, Hemorrhage, Other Specified Gastrointestinal Disorders *	82,250	5.9	0.053	0.014	0.000	1.054	1.025	1.083
HCC365_40	HCC40 Rheumatoid Arthritis and Inflammatory Connective Tissue Disease	81,912	5.9	0.086	0.014	<.0001	1.087	1.057	1.117
HCC365_46	HCC46 Severe Hematological Disorders	13,156	1.0	0.123	0.030	<.0001	1.131	1.065	1.201
HCC365_47	HCC47 Disorders of Immunity	33,676	2.4	0.062	0.020	0.002	1.064	1.022	1.107
HCC365_49	HCC49 Iron Deficiency and Other/Unspecified Anemias and Blood Disease	602,721	43.6	0.051	0.008	<.0001	1.053	1.037	1.069
HCC365_64	HCC64 Profound Mental Retardation/Developmental Disability	647	0.0	0.471	0.127	0.000	1.623	1.259	2.092
HCC365_65	HCC65 Severe Mental Retardation/Developmental Disability	1,199	0.1	0.312	0.103	0.003	1.373	1.116	1.689
HCC365_66	HCC66 Moderate Mental Retardation/Developmental Disability	871	0.1	0.134	0.136	0.326	1.139	0.867	1.496

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
HCC365_70	HCC70 Quadriplegia	6,450	0.5	0.152	0.043	0.000	1.167	1.071	1.272
HCC365_71	HCC71 Paraplegia	7,860	0.6	0.095	0.042	0.022	1.097	1.010	1.193
ALS_MDys_MS	HCC73 Amyotrophic Lateral Sclerosis and Other Motor Neuron Disease, HCC76 Muscular Dystrophy, HCC77 Multiple Sclerosis	11,562	0.8	0.106	0.038	0.005	1.106	1.026	1.193
HCC365_75	HCC75 Polyneuropathy	145,921	10.6	0.031	0.012	0.008	1.030	1.006	1.055
HCC365_78	HCC78 Parkinson's and Huntington's Diseases	51,947	3.8	0.037	0.018	0.039	1.038	1.001	1.076
HCC_prox_82	HCC82 Respirator Dependence/Tracheostomy Status *	4,714	0.3	0.137	0.047	0.004	1.145	1.042	1.258
HCC_prox_83	HCC83 Respiratory Arrest *	466	0.0	0.148	0.161	0.359	1.157	0.838	1.597
HCC365_84	HCC84 Cardio-Respiratory Failure and Shock	211,849	15.3	0.054	0.009	<.0001	1.053	1.034	1.073
HCC365_85	HCC85 Congestive Heart Failure	468,619	33.9	0.222	0.009	<.0001	1.249	1.227	1.270
HCC_prox_86	HCC86 Acute Myocardial Infarction *	21,630	1.6	0.072	0.025	0.004	1.072	1.020	1.127
Angine_Unst_Pector	HCC87 Unstable Angina and Other Acute Ischemic Heart Disease, HCC88 Angina Pectoris *	24,659	1.8	0.042	0.025	0.092	1.040	0.989	1.093
HCC_prox_89	HCC89 Coronary Atherosclerosis/Other Chronic Ischemic Heart Disease *	398,854	28.8	0.056	0.008	<.0001	1.059	1.042	1.076
HCC_prox_90	HCC90 Heart Infection/Inflammation, Except Rheumatic *	7,696	0.6	0.059	0.040	0.143	1.060	0.977	1.149
HCC365_91	HCC91 Valvular and Rheumatic Heart Disease	194,409	14.1	0.049	0.010	<.0001	1.048	1.028	1.068
HCC365_96	HCC96 Specified Heart Arrhythmias	441,435	31.9	0.141	0.008	<.0001	1.149	1.131	1.167
HCC365_109	HCC109 Other Circulatory Disease	181,262	13.1	0.019	0.010	0.049	1.016	0.997	1.036
HCC365_111	HCC111 Chronic Obstructive Pulmonary Disease	372,930	27.0	0.252	0.008	<.0001	1.288	1.267	1.310
HCC365_112	HCC112 Fibrosis of Lung and Other Chronic Lung Disorders	19,068	1.4	0.165	0.027	<.0001	1.177	1.114	1.244
HCC365_113	HCC113 Asthma	50,036	3.6	0.085	0.020	<.0001	1.086	1.043	1.132
HCC365_114	HCC114 Aspiration and Specified Bacterial Pneumonias	77,164	5.6	0.118	0.014	<.0001	1.125	1.095	1.156
HCC365_116	HCC116 Viral and Unspecified Pneumonia, Pleurisy	199,991	14.5	0.114	0.009	<.0001	1.122	1.102	1.143

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

Variable Name in Model	Covariate	Count	Percent Total	Estimate	Std. Error	P value	Odds Ratios	OR 95% Lower CL	OR 95% Upper CL
HCC_prox_117	HCC117 Pleural Effusion/Pneumothorax	47,744	3.5	0.046	0.017	0.006	1.046	1.011	1.082
HCC365_132	HCC132 Kidney Transplant Status	4,825	0.3	0.284	0.049	<.0001	1.321	1.197	1.459
HCC365_134	HCC134 Dialysis Status	32,886	2.4	0.039	0.033	0.229	1.036	0.970	1.106
Acute_Unspecified_Re	HCC135 Acute Renal Failure, HCC140 Unspecified Renal Failure	345,152	25.0	0.200	0.009	<.0001	1.218	1.196	1.239
AdvCKD	HCC136 Chronic Kidney Disease, Stage 5, HCC137 Chronic Kidney Disease, Severe (Stage 4), HCC138 Chronic Kidney Disease, Moderate (Stage 3), HCC139 Chronic Kidney Disease, Mild or Unspecified (Stages 1-2 or Unspecified)	149,312	10.8	0.184	0.012	<.0001	1.197	1.170	1.225
HCC_prox_141	HCC141 Nephritis *	2,045	0.1	0.102	0.081	0.208	1.106	0.940	1.302
HCC_prox_142	HCC142 Urinary Obstruction and Retention *	108,464	7.8	0.070	0.013	<.0001	1.070	1.043	1.098
HCC365_144	HCC144 Urinary Tract Infection	408,285	29.5	0.088	0.008	<.0001	1.094	1.076	1.113
HCC_prox_145	HCC145 Other Urinary Tract Disorders *	74,122	5.4	0.054	0.015	0.000	1.055	1.024	1.086
HCC_prox_157	HCC157 Pressure Ulcer of Skin with Necrosis Through to Muscle, Tendon, or Bone *	5,794	0.4	0.341	0.043	<.0001	1.410	1.292	1.538
HCC_prox_158	HCC158 Pressure Ulcer of Skin with Full Thickness Skin Loss *	12,723	0.9	0.217	0.029	<.0001	1.242	1.171	1.317
HCC_prox_159	HCC159 Pressure Ulcer of Skin with Partial Thickness Skin Loss *	20,936	1.5	0.114	0.024	<.0001	1.121	1.068	1.177
HCC_prox_160	HCC160 Pressure Pre-Ulcer Skin Changes or Unspecified Stage *	20,166	1.5	0.083	0.025	0.001	1.086	1.033	1.142
HCC365_161	HCC161 Chronic Ulcer of Skin, Except Pressure	42,025	3.0	0.131	0.018	<.0001	1.138	1.099	1.180
HCC365_164	HCC164 Cellulitis, Local Skin Infection	121,210	8.8	0.082	0.012	<.0001	1.087	1.062	1.113
HCC_prox_165	HCC165 Other Dermatological Disorders *	39,142	2.8	0.022	0.021	0.288	1.018	0.977	1.061
HCC_prox_169	HCC169 Vertebral Fractures without Spinal Cord Injury *	23,554	1.7	0.018	0.029	0.529	1.017	0.960	1.077
HCC365_186	HCC186 Major Organ Transplant or Replacement Status	2,838	0.2	0.107	0.063	0.091	1.112	0.980	1.262

Table 2-8. Potentially Preventable Unplanned Readmission Measure for 30 Days Post Discharge from Skilled Nursing Facilities: Logistic Regression Model Results in 2013 (continued)

			Percent		Std.		Odds	OR 95% Lower	OR 95%
Variable Name in Model	Covariate	Count	Total	Estimate	Error	P value	Ratios	CL	Upper CL
HCC365_188	HCC188 Artificial Openings for Feeding or Elimination	37,705	2.7	0.138	0.018	<.0001	1.147	1.106	1.190
HCC365_190	HCC190 Amputation Status, Upper Limb	2,066	0.1	0.118	0.077	0.125	1.130	0.969	1.318
	Original Reason for Entitlement								
origDisabled	Original reason for entitlement is disabled	289,851	21.0	0.078	0.011	<.0001	1.083	1.060	1.106
	Prior Acute ICU/CCU Days (Ref: p_ICU_CCU_0)								
anyICUdays	At least one day in prior proximal ICU (INTNSV_CARE_DAY_CNT_prox)	374,166	27.1	0.012	0.008	0.151	1.014	0.997	1.031
	Prior Acute Care Length of Stay (Reference group: LOS 1-3 Days)								
los4_7	LOS btwn 4 & 7 days (based on proximal IPPS)	638,548	46.2	0.106	0.009	<.0001	1.116	1.095	1.137
los8_14	LOS btwn 8 & 14 days (based on proximal IPPS)	254,889	18.4	0.188	0.011	<.0001	1.214	1.187	1.242
los15_GT	LOS GT 14 days (based on proximal IPPS)	81,567	5.9	0.194	0.017	<.0001	1.225	1.184	1.267
	Prior Acute Care Utilization-Count of prior stays								
prior_1_3	1-3 IPPS stays in 365 days prior	633,094	45.8	0.195	0.009	<.0001	1.222	1.199	1.244
prior_4_6	4-6 IPPS stays in 365 days prior	84,176	6.1	0.496	0.015	<.0001	1.662	1.613	1.712
prior_7_9	7-9 IPPS stays in 365 days prior	13,754	1.0	0.747	0.025	<.0001	2.144	2.038	2.255
prior_GE_10	10+ IPPS stays in 365 days prior	3,793	0.3	1.049	0.041	<.0001	2.905	2.679	3.151

Note: Number of Observations in 2013: 1,382,654. The C-Statistic was 0.71.

Source: RTI International analysis of Medicare claims data, 2013. (RTI program reference: idxSNF05\_018b.xls)

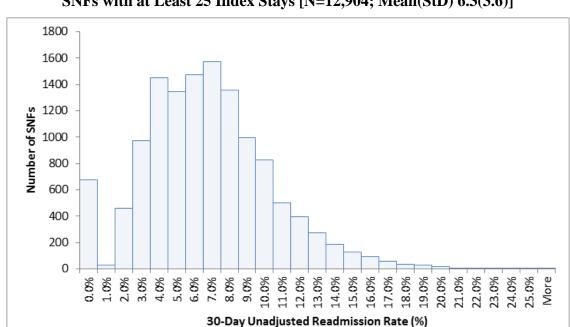


Figure 2-2. Distribution of Unadjusted Potentially Preventable Readmission Rates among SNFs with at Least 25 Index Stays [N=12,904; Mean(StD) 6.3(3.6)]

Source: RTI International analysis of Medicare claims data, 2013. (RTI program reference: idxSNF05\_017b.xls)

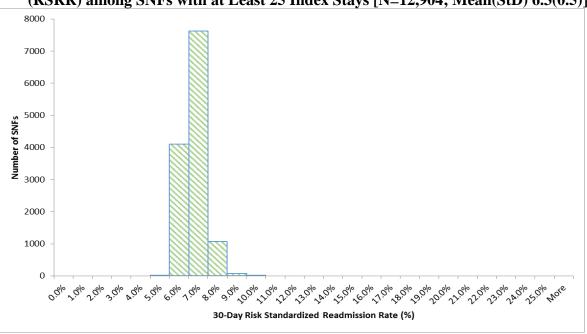


Figure 2-3. Distribution of Risk Standardized Potentially Preventable Readmission Rates (RSRR) among SNFs with at Least 25 Index Stavs [N=12,904; Mean(StD) 6.3(0.5)]

Source: RTI International analysis of Medicare claims data, 2013. (RTI program reference: idxSNF05\_017b.xls) Note: Scales differ.

## **APPENDIX 3**

## DRUG REGIMEN REVIEW CONDUCTED WITH FOLLOW-UP FOR IDENTIFIED ISSUES- POST ACUTE CARE (PAC) SKILLED NURSING FACILITY (SNF) QUALITY REPORTING PROGRAM (QRP)

Table 1 below summarizes the setting specific language used to describe the resident or patient within the PAC setting. There are no other differences in the content language within each Drug Regimen Review quality measure item.

Table 3-1
Drug Regimen Review Quality Measure Setting-Specific Language

SNF	IRF	LTCH
Beginning of stay	Beginning of stay	Beginning of stay
N2001 Drug Regimen Review:	N. 2001 Drug Regimen Review:	N. 2001 Drug Regimen Review:
Did a complete drug regimen review identify potential clinically significant medication issues?	Did a complete drug regimen review identify potential clinically significant medication issues?	Did a complete drug regimen review identify potential clinically significant medication issues?
☐ 0 - No - No issues found during review	0 - No - No issues found during review	☐ 0 - No - No issues found during review
1 - Yes - Issues found during review	1 - Yes - Issues found during review	1 - Yes - Issues found during review
☐ 9 - NA – Resident is not taking any medications	☐ 9 - NA – Patient is not taking any medications	☐ 9 - NA – <mark>Patient</mark> is not taking any medications
N. 2003 Medication Follow-up: Did the facility contact a physician (or physician-designee) by midnight of the next calendar day and complete prescribed/recommended actions in response to the identified potential clinically significant medication issues?  □ 0 - No □ 1 - Yes	N. 2003 Medication Follow-up: Did the facility contact a physician (or physician-designee) by midnight of the next calendar day and complete prescribed/recommended actions in response to the identified potential clinically significant medication issues?  □ 0 - No □ 1 - Yes	N. 2003 Medication Follow-up: Did the facility contact a physician (or physician-designee) by midnight of the next calendar day and complete prescribed/recommended actions in response to the identified potential clinically significant medication issues?  □ 0 - No □ 1 - Yes

Table 3-1
Drug Regimen Review Quality Measure Setting-Specific Language (continued)

SNF	IRF	LTCH
End of stay	End of stay	End of stay
N. 2005 Medication Intervention:	N. 2005 Medication Intervention:	N. 2005 Medication Intervention:
Did the facility contact and complete physician (or physician-designee) prescribed/recommended actions by midnight of the next calendar day each time potential clinically significant medication issues were identified since the Admission?	Did the facility contact and complete physician (or physician-designee) prescribed/recommended actions by midnight of the next calendar day each time potential clinically significant medication issues were identified since the Admission?	Did the facility contact and complete physician (or physician-designee) prescribed/recommended actions by midnight of the next calendar day each time potential clinically significant medication issues were identified since the Admission?
□ 0 - No	□ 0 - No	□ 0 - No
□ 1 - Yes	☐ 1 - Yes	☐ 1 - Yes
□ 9 - NA -There were no potential clinically significant medication issues identified since Admission or resident is not taking any medications.	□ 9 - NA -There were no potential clinically significant medication issues identified since Admission or patient is not taking any medications.	□ 9 - NA -There were no potential clinically significant medication issues identified since Admission or patient is not taking any medications.