

Fall 2024 Updates

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Resident Assessment Instrument version 1.19.1 October 2024

Civil Money Penalties (CMP)
fund is currently over
33 million dollars!

Please visit
NC Culture Change Coalition
for ideas and apply for
funding for your facilities

<https://www.ncculturechangecoalition.org/>





QIN-QIO

Quality Innovation Network -

Quality Improvement Organizations

CENTERS FOR MEDICARE & MEDICAID SERVICES

QUALITY IMPROVEMENT & INNOVATION GROUP

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No cost resources and services for facilities

Non-regulatory



CENTER OF
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IN NURSING FACILITIES

nursinghomebehavioralhealth.org

Disclaimer:

This presentation is not a substitute for reading and reviewing the

- ❖ Long-Term Care Resident Assessment Instrument 3.0 User's Manual Version 1.19.1, October 2024
 - ❖ MDS Item Sets Version 1.19.1, October 2024
- or
- ❖ State Operations Manual Appendix PP, Revised 8/8/24

Purpose and Objectives to Review:

- RAI and MDS changes starting October 1, 2024
- CMS Schizophrenia Audits
- PASRR Information
- Most Frequent Citations Related to MDS and the Regulations
- Helpful Reports in iQIES
- Resources

A2121 Provision of Current Reconciled Medication List to Subsequent Provider at Discharge

- Removed examples of coding that involved remaining in the nursing home or SNF or LTC
- If the resident remains long term care after the SNF coverage ends, code 1: Yes, that you provided the medication list to the subsequent provider, even if that provider is you.

RAI page A-44

A2121. Provision of Current Reconciled Medication List to Subsequent Provider at Discharge

Complete only if A0310H = 1 and A2105 = 02-12

Enter Code

At the time of discharge to another provider, did your facility provide the resident's current reconciled medication list to the subsequent provider?

0. **No** - Current reconciled medication list not provided to the subsequent provider → Skip to A2200, Previous Assessment Reference Date for Significant Correction
1. **Yes** - Current reconciled medication list provided to the subsequent provider

Brief Interview for Mental Status

- Coding Tips:
- If all the BIMS items are coded with a dash (-), then C0500, BIMS Summary Score must also be coded with a dash (-).

RAI page C-17

Section GG

- Now called “Functional Abilities”
- All Goals for the PPS 5-day and references to goals have been removed
- The Goals column for the PPS 5-day is gone.
- All references to the Goals column have been removed.
- Goals are no longer a SNF QRP item

GG0310I, Personal Hygiene

- Personal hygiene involves the ability to maintain personal hygiene, including combing hair, shaving, applying makeup, and washing and drying face and hands (excludes baths, showers, and oral hygiene)
- Examples have been adjusted

GG0170 Mobility

- **Coding Tips for GG0170M, 1 step (curb); GG0170N, 4 steps; and GG0170O, 12 steps**
- If, at the time of the assessment, a resident is unable to complete the activity because of a physician-prescribed restriction *of no stair climbing, they may be able to complete the stair activities safely by some other means (e.g., stair lift, bumping/scooting on their buttocks). If so, code based on the type and amount of assistance required to complete the activity. If, at the time of assessment, a resident is unable to complete the stair activities because of a physician-prescribed bedrest, code the stair activity using the appropriate “activity not attempted” code.*
- *While a resident may take a break between ascending or descending the 4 steps or 12 steps, once they start the activity, they must be able to ascend (or descend) all the steps, by any safe means, without taking more than a brief rest break to consider the stair activity completed.*

H0100: Appliances

- **DEFINITIONS**
EXTERNAL CATHETER

Device attached to the shaft of the penis like a condom, *a female external catheter, or other non-invasive urine output management device or system that routes urine* to a drainage bag.

- **Coding Tips and Special Populations**

Female external catheters and other non-invasive urine output management devices or systems should be coded as external catheters (H0100B).

RAI pages H-2 and H-3

Clarification

- Item I2100 Septicemia:
- For sepsis to be considered septicemia, there needs to be inflammation due to sepsis and evidence of a microbial process.
- If the medical record reflects inflammation due to sepsis and evidence of a microbial process, code I2100, Septicemia.
- If the medical record does not reflect inflammation due to sepsis and evidence of a microbial process, enter the sepsis diagnosis and ICD code in item I8000, Additional Active Diagnoses.

K0520 Modified Definition of Feeding Tube

DEFINITIONS

FEEDING TUBE

Presence of any type of tube that can deliver food/ nutritional substances/ fluids directly into the gastrointestinal system. Examples include, but are not limited to, nasogastric tubes, gastrostomy tubes, jejunostomy tubes, percutaneous endoscopic gastrostomy (PEG) tubes. RAI page K-10

- Coding tip matches the black box definition
- **Coding Tip for K0520B**
- Only feeding tubes that are used to deliver nutritive substances and/or hydration during the assessment period are coded in K0520B. RAI page K-12

K0520 Parenteral/IV Feeding

- **Coding Tips for K0520A**

- IV fluids can be coded in K0520A if needed to prevent dehydration if the additional fluid intake is specifically needed for nutrition and/or hydration. Prevention of dehydration should be clinically indicated and supporting documentation should be provided in the medical record. RAI page K-12

N0415K, Anticonvulsant

- **N0415K1. Anticonvulsant:** *Check if an anticonvulsant medication was taken by the resident at any time during the 7-day observation period (or since admission/entry or reentry if less than 7 days).*
- **N0415K2. Anticonvulsant:** *Check if there is an indication noted for all anticonvulsant medications taken by the resident any time during the observation period (or since admission/entry or reentry if less than 7 days).* RAI page N-8
- Anticonvulsants are used in the treatment of epileptic seizures, and neuropathic pain. Also, for bipolar disorder and borderline personality disorder since many seem to act as mood stabilizers.

0011001, IV Access

- **0011001, IV Access** Code IV access, which refers to a catheter inserted into a vein for a variety of clinical reasons, including long-term medication administration, large volumes of blood or fluid, frequent access for blood samples, intravenous fluid administration, total parenteral nutrition (TPN), or, in some instances, the measurement of central venous pressure. *An arteriovenous (AV) fistula does not meet the definition of IV Access for 0011001.*

RAI page O-9

- An A-V fistula is a surgical connection made between an artery and a vein, created by a vascular specialist.

Vaccinations: from the LTC Survey Pathway

- **Influenza, Pneumococcal, and COVID-19 Immunizations for Residents:**
- Review the records for documentation of: Screening and eligibility to receive the vaccine(s);
- The provision of education related to the influenza, pneumococcal, and COVID-19 vaccines;
- The administration of vaccines in accordance with national recommendations, which includes doses administered.
- Facilities must follow the CDC and Advisory Committee on Immunization Practices (ACIP) recommendations for vaccines; and
- Allowing a resident or representative to accept or refuse the influenza, pneumococcal, and COVID-19 vaccines. If not provided, documentation as to why the vaccine(s) was not provided.

Vaccinations: from the LTC Survey Pathway

- For surveys occurring during influenza season, unavailability of the influenza vaccine can be a valid reason why a facility has not implemented the influenza vaccine program, especially during the early weeks of the influenza season. Similarly, pneumococcal or COVID-19 vaccine supplies may be limited anytime of the year. Ask the facility to demonstrate that: The vaccine has been ordered and the facility received a confirmation of the order indicating that the vaccine has been shipped or that the product is not available but will be shipped when the supply is available; and
- Plans are developed on how and when the vaccines will be administered when they are available.
- As necessary, determine if the facility developed influenza, pneumococcal, and COVID-19 vaccine policies and procedures for residents....
- Survey Resources: <https://www.cms.gov/Medicare/Provider-Enrollment-and-Certification/GuidanceforLawsAndRegulations/Nursing-Homes>

O0300: Pneumococcal Vaccine Examples

- Pneumococcal Vaccine examples now include PCV20
- PCV13, PCV20, PPSV23
- RAI starting on page O-17

Pneumococcal Vaccine

- If a resident has received one or more pneumococcal vaccinations and is indicated to get an additional pneumococcal vaccination but is not yet eligible for the next vaccination because the recommended time interval between vaccines has not lapsed, O0300A is coded 1, yes, indicating the resident's pneumococcal vaccination is up to date.
- Advisory Committee on Immunization Practices (ACIP) Vaccine Recommendations and Guidelines
<https://www.cdc.gov/vaccines/hcp/acip-recs/index.html>

O0350: Resident's COVID-19 vaccination is up to date

O0350. Resident's COVID-19 vaccination is up to date

Enter Code

0. No, resident is not up to date
1. Yes, resident is up to date

• Steps for Assessment

- *Vaccination status may be determined based on information from any available source. Review the resident's medical record or documentation of COVID-19 vaccination and/or interview the resident, family or other caregivers or healthcare providers to determine whether the resident is up to date with their COVID-19 vaccine.*
- *If the resident is **not up to date**, and the facility has the vaccine available, ask the resident if they would like to receive the COVID-19 vaccine.*

RAI starting on page O-19

If you don't know the status= not up to date

O0350: Resident's COVID-19 (continued)

- **Coding Instructions**

- *Code 0, No, resident is not up to date if the resident does not meet the CDC's definition of up to date. This includes residents who have not received one or more recommended COVID-19 vaccine doses **for any reason** including medical, religious, or other qualified exemptions.*
- *This includes residents for whom vaccination status cannot be determined.*
- *Code 1, Yes, resident is up to date if the resident meets the CDC's definition of up to date.*
- *A dash is a valid response, indicating the item was not assessed. CMS expects dash use to be a rare occurrence.*

- **Coding Tip**

- *Current COVID-19 vaccine recommendations are available on the Centers for Disease Control and Prevention's (CDC's) webpage "Stay Up to Date with COVID-19 Vaccines" at <https://www.cdc.gov/coronavirus/2019-ncov/vaccines/stay-up-to-date.html>*

The Cross-Over Rule, Chapter 5.7

- *When item sets are updated, a situation may exist that will prevent providers from correcting the target date of any assessment crossing over from October 1 of a given year. That is, providers may not submit a modification to change a target date on an assessment completed prior to October 1 of a given year to a target date on or after October 1 of the same year, nor can they submit a modification to change a target date on an assessment completed on or after October 1 of a given year to a target date prior to October 1 of a given year when the MDS item sets have had substantial changes.*
- *When the MDS item sets have had significant changes, including the omission and addition of many items or significant changes to existing items, clinicians will be required to collect and code new items, may have different look-back periods, or may need to code the MDS according to changes in the coding requirements.* It is the target date of the assessment that identifies the required version of the item set, and, because of the substantial changes *that may exist between versions of* the item sets, they are not interchangeable. Therefore, *commonly when there are updates to item sets,* providers may not change target dates on assessments crossing over October 1 *of specific years.* RAI page 5-12

5.8 Special Manual Record Correction Request

- A few types of errors in a record in iQIES cannot be corrected with an automated Modification or Inactivation request. These errors are:
 - The record has the wrong unit certification or licensure designation in Item A0410.
 - The record has the wrong state code or facility ID in the control Items STATE_CD or FAC_ID.
 - *The record submitted was not for OBRA or Medicare Part A purposes.*
 - The record is a test record inadvertently submitted as production.

Chapter 5 Submission and Correction

- *When a facility erroneously submits a record that was not for OBRA or Medicare Part A purposes, CMS does not have the authority to collect the data contained in the record. An inactivation request will not fix the problem, since it will leave the erroneously submitted record in the history file, that is, the CMS database. A manual deletion is necessary to completely remove the erroneously submitted record and associated information from the CMS database.*
- *In instances in which an erroneous PPS assessment is combined with an OBRA-required assessment, if the item set code does not change, then a modification can be completed. If the item set code does change as a result of a modification, the provider must complete an MDS 3.0 Manual Assessment Correction/Deletion Request. This action will completely remove the assessment from the database. As indicated, the provider would complete and submit a new, stand-alone OBRA assessment.*

Manual Deletion Form

NOTE: Assessment item errors, other than those listed below, must be corrected and resubmitted using Correction Policy procedures.

Please Type or Print Legibly
All Fields are Required

Delete Test Record Correct A0410 Value Delete Wrong FAC_ID Not CMS Required***

Facility Information

Facility Name: ID (FAC_ID):

Requestor (Administrator/Owner) Information

Name (full name): Title:
E-mail Address: Phone Number:

Resident Information

First Name: Last Name:
SSN: Birth Date: Gender:
Resident ID:*

Record Information

A0310A Value: A0310B Value: A0310C Value: A0310D Value: A0310F Value:
Target Date:** Assessment ID:*

Submission Information

Submission Date: Submission ID:*

A0410 (Submission Requirement) Values

Submitted (Incorrect) Value: **3** Correct Value:

* RES_INT_ID, ASMT_ID, and SUBMISSION ID are found on the Final Validation Report

** Target Date is:
MDS Item A2300 (Assessment Reference Date) for an assessment record
MDS Item A2000 (Discharge Date) for a discharge record
MDS Item A1600 (Entry Date) for a reentry record
*** Record is not for OBRA and not for Medicare Part A PPS

Submit completed and signed form to the iQIES Service Center by **Certified Mail** through the US Postal Service.

GDIT
iQIES Service Center
4800 Westown Pkwy, Suite 360
West Des Moines, IA 50266

Signature - Administrator or Owner (Please circle one) _____ Date _____
Submit completed and signed form to your State Agency via **Certified Mail** through the US Postal Service. Your State Agency will approve, sign, and forward your request to the iQIES Service Center.

Signature - State Agency Authorizer _____ Date _____
The request must be sent **Certified Mail** through the US Postal Service.

All requests require State Agency authorization.

Forms forwarded to the iQIES Service Center without a State Agency signature will be rejected.

iQIES Service Center - Internal Use:

<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
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- Nearly all of the information required to complete this form can be found on the Final Validation Report:
- Facility ID
- Resident ID
- Record Information
- Submission Information

Any questions call 919-909-9256

Mail to:

Janet Brooks
RAI Education Coordinator
Division of Health Service Regulation
Nursing Home Licensure & Certification Section
2711 Mail Service Center
Raleigh, NC 27699

A0410

A0410. Unit Certification or Licensure Designation

Enter Code

1. Unit is neither Medicare nor Medicaid certified and MDS data is not required by the State
2. Unit is neither Medicare nor Medicaid certified but MDS data is required by the State
3. Unit is Medicare and/or Medicaid certified

- A0410 Unit Certification or Licensure Designation

1. Unit is neither Medicare nor Medicaid certified and MDS data is not required by the State

2. Unit is neither Medicare nor Medicaid certified but MDS data is required by the State

3. Unit is Medicare and/or Medicaid certified

Should always be '3' unless it should not be transmitted then '1' would be coded

2.14 Determining the Item Set for an MDS Record

- The item set for a particular MDS record is completely determined by the reason for assessment items (A0310A, A0310B, A0310F, and A0310H)....
- The first lookup table is for nursing home records. The first 4 columns are entries for the reason for assessment (RFA) items A0310A, A0310B, A0310F, and A0310H. To determine the item set for a record, locate the row that includes the values of items A0310A, A0310B, A0310F, and A0310H for that record. When the row is located, then the item set is identified in the item set code (ISC) and Description columns for that row. If the combination of items A0310A, A0310B, A0310F, and A0310H values for the record cannot be located in any row, then that combination of RFAs is not allowed and will be rejected by iQIES.

RAI page 2-57

Nursing Home Item Set Code (ISC) Reference Table

OBRA RFA (A0310A)	PPS RFA (A0310B)	Entry/ Discharge (A0310F)	Part A PPS Discharge (A0310H)	ISC	Description
01, 03, 04, 05	01, 99	10, 11, 99	0, 1	NC	Comprehensive
02, 06	01, 99	10, 11, 99	0, 1	NQ	Quarterly
99	01	10, 11, 99	0, 1	NP	PPS
99	08	99	0	IPA	PPS (Optional)
99	99	10, 11	0, 1	ND	OBRA Discharge
99	99	01, 12	0	NT	Tracking
99	99	99	1	NPE	Part A PPS Discharge

QSO-23-05-NH

Updates to the Nursing Home Care Compare Website and Five Star Quality Rating System:

Adjusting Quality Measure Ratings Based on Erroneous Schizophrenia Coding, and Posting Citations Under Dispute

<https://www.cms.gov/files/document/qso-23-05-nh.pdf>

1/18/2023

16000 Schizophrenia

Memorandum Summary Adjusting Quality Measure Ratings:

CMS will be conducting audits of schizophrenia coding in the Minimum Data Set data and based upon the results, adjust the Nursing Home Care Compare quality measure star ratings for facilities whose audits reveal inaccurate coding.

CMS is concerned that some nursing homes have erroneously coded residents as having schizophrenia, which can mask the facilities' true rate of antipsychotic medication use.

Facilities that have coding inaccuracies identified through the schizophrenia MDS audit will have their QM ratings adjusted as follows:

- The Overall QM and long stay QM ratings will be downgraded to one star for six months (this drops the facility's overall star rating by one star).
- The short stay QM rating will be suppressed for six months.
- The long stay antipsychotic QM will be suppressed for 12 months.

Determinations:

1. Sufficient documentation of behaviors was not present prior to establishing or validating the diagnosis of schizophrenia.
2. Residents were given a new diagnosis of schizophrenia without comprehensive medical evaluations, excluding other medical conditions.
3. Residents were given a new diagnosis of schizophrenia without comprehensive psychiatric evaluations.
4. Sufficient documentation was not present to support a 'history of schizophrenia' as noted in the medical record.
5. Sufficient documentation of comprehensive evaluations that validate the accuracy of the diagnosis of schizophrenia on-admission, as noted in the medical record, was not present.
6. Coding of schizophrenia on MDS assessments reflect inaccuracies compared to the reviewed medical records.
7. Coding of antipsychotic medications on MDS assessments reflect inaccuracies compared to the reviewed medical records.
8. Sufficient documentation was not present to support the removal of schizophrenia as an active diagnosis.
9. The reviewed medical records lacked the required MDS assessment modifications to remove an inaccurate diagnosis of schizophrenia.

SCHIZOPHRENIA IN NURSING FACILITIES: Validating Diagnosis and Planning for Appropriate Care

Schizophrenia is a serious life-long brain disorder with a wide range of symptoms that affect a person's thoughts, emotions, and behaviors. A new diagnosis after the age of 40 is rare and after age 65 is uncommon. A new onset of Schizophrenia in a post-acute and long term care setting should not be coded on the Minimum Data Set (MDS) 3.0 Resident Assessment Instrument (RAI) Manual (Item I6000) unless there is documentation of the diagnosis by a qualified clinician, using evidence-based criteria and professional standards, such as the Diagnostic and Statistical Manual of Mental Disorders Fifth Edition Revision (DSM-5-TR).

Checkpoints to verify the accuracy of a Schizophrenia diagnosis.

- Documentation should include supporting evidence of a history of symptoms beginning at an appropriate age. Schizophrenia symptoms typically become apparent at ages 16-30.
- Ensure dementia, other mental health disorders, or medical conditions with similar symptoms have been ruled out.
- Collaborate with the resident and family to obtain symptoms, psychiatric and family history.
- Efforts should be made to obtain records from identified outpatient behavioral health provider to confirm the diagnosis.
- Further assessment and review of the resident's history may be necessary if the diagnosis is based only on hospital records. Involve the appropriate medical clinicians in the nursing facility to ensure an appropriate diagnosis.
- If an antipsychotic is added during the hospital admission, verify and assess the need for continued use. A diagnosis of Schizophrenia should not be added to solely support the use of an antipsychotic medication.

Preadmission Screening and Resident Review (PASRR):

Many residents with a Schizophrenia diagnosis meet the criteria for a Level II PASRR. Consider these questions:

- Is proper PASRR Level I documentation present?
- If Level II was required, was it completed?
- Are any specialized services recommended in the PASRR review?
- Are specialized services being provided?
- Is the PASRR status properly documented in MDS Item # A1500, A1510?



Individualized Care Planning:

- Include the resident and family in the care planning process.
- Ensure the care plan interventions are individualized.
- Obtain recommendations from the resident and family for effective symptom management strategies.
- If interventions are not effective, they should be discontinued, and new interventions implemented.
- Incorporate the PASRR review details into the care plan.
- Document in the medical record the use and effectiveness of all pharmacological and nonpharmacological interventions.
- When a pharmacological intervention is instituted, document the effectiveness and attempts made to use the lowest effective dose under the supervision of a qualified medical clinician in collaboration with the consultant pharmacist. *See §483.45(c)(3) and (e), F758 Psychotropic drugs*

Potential Appropriate Interventions Include:

- Individual therapy
- Group therapy
- Individualized, social, and psychosocial activities
- Train nursing facility staff to understand the resident's condition and appropriate interventions.

A1500 –Preadmission Screening and Resident Review (PASRR)

- PASRR is a preadmission screening process.
- A positive screen indicates the resident has a mental illness, intellectual disability, or a related condition.
- A1500 documents whether a PASRR Level II determination has been issued.
- Reports on the results of the PASRR process.
- A1500 is only completed on the OBRA comprehensive MDS assessments.

Pre-Admission Screening and Resident Review

- Not everyone with MI has a Level II PASRR determination.
- Everyone with ID/DD should have a Level II PASRR determination.

Further Information and Resources

- RAI page 2-30 through 2-31
- F644, F645, F646
- PASRR Help Desk 888-245-0179, 919-813-5603

<https://medicaid.ncdhhs.gov/providers/programs-and-services/long-term-care/pre-admission-screening-and-resident-review-pasrr>

<https://ncliftss.acentra.com/pasrr/>

Pre-Admission Screening and Resident Review

- **When is PASRR required?**
- PASRR is required anytime someone is being admitted into a Medicaid-certified nursing facility. The PASRR determination should be recent.
 - All individuals who are admitted to a Medicaid certified nursing facility, must have a Level I PASRR completed to screen for possible mental illness (MI), intellectual disability (ID), (“mental retardation” (MR) in federal regulation)/developmental disability (DD), or related conditions. F645
- Before the end of a time-limited approval if a person is expected to remain in the nursing home for longer than the approved time.
- Any Level II resident experiencing a significant change in status.
- Any Level I resident who experience a psychiatric episode, have a new psychiatric diagnosis or have been placed on antipsychotic medications should have a Level II PASRR referral made.

Halted PASRR

Halted – Level II Authorization No end date, no restrictions.

(indicates Dementia primary or Does Not Meet Level II Target Population Criteria)

Halted – Level II authorizations halted due to dementia primary, terminal prognosis, or does not meet Level II Target Population Criteria after further assessment. No restrictions, no end date unless a change in condition.

1/19/24

<https://medicaid.ncdhhs.gov/documents/providers/programs-services/pasrr/pasrr-authorizations-quick-reference/download>

North Carolina PASRR: Skilled Nursing Facility

Authorization Codes & Corresponding Time Frames/ Restrictions FOLLOWING CODES NOT VALID FOR Adult Care Home Admission or Placement	
A	No end date no mental or behavioral health restrictions.
H	Halted – Level II Authorization no end date, no restrictions. (Indicates Dementia primary or Does Not Meet Level II Target Population Criteria)
B	Level II: No end date, no limitation unless change in condition. No specialized services required.
C	Level II: No end date, no limitation unless change in condition. Specialized services required.
E	Level II: 30-Day Rehabilitation Services Authorization only.
D	Level II: 7-Day Respite or Emergency Placement Authorization only.
J	Level II: 1-year Authorization for placement at a <u>Locked</u> State Psychiatric Hospital or <u>State Operated</u> Nursing Facility only.
F	Level II: 30, 60 or 90 Day Authorization for Time Limited Skilled Nursing Facility stays
Z	Level II: Denial. Nursing facility placement is <u>not</u> appropriate.

For additional information and/or clarification please review the nursing facility clinical coverage policy ZB1 located at:
<https://amu.ncdhhs.gov/document/facility-services-clinical-coverage-policies>

North Carolina Skilled Nursing Facility Authorization Codes & Corresponding Time Frames/Restrictions

A	No end date unless change in condition no mental or IDD diagnoses identified
H	Halted – Level II authorizations halted due to dementia primary, terminal prognosis, or does not meet Level II Target Population Criteria after further assessment. No restrictions, no end date unless a change in condition.
B	Level II: no end date, no limitation unless change in condition, No Specialized services required.
C	Level II: no end date, no limitation unless change in condition, Specialized services required.
E	Level II: 30-Day Rehabilitation Services Authorization Only
D	Level II: 7-Day Respite or Emergency Placement Authorization Only.
J	Level II: 1-year Authorization for placement at a <u>Locked</u> State Psychiatric Hospital or <u>State Operated</u> Nursing Facility Only
F	Level II: 30, 60 or 90-Day Authorization for Time limited Skilled Nursing Facility Stays
Z	Level II: Denial. Nursing facility placement is NOT appropriate

Halted Level II PASRR

- Halted – Level II authorizations halted due to dementia primary, terminal prognosis, or does not meet Level II Target Population Criteria after further assessment. No restrictions, no end date unless a change in condition
 - *Continues to be a level II PASRR*
 - *Should your facility say the H designation PASRR is a Level I, the burden of proof is on the facility to provide that documentation from NCMUST*
 - *If the H designation is related to dementia, the physician must document dementia is the primary diagnosis*
 - *PASRR determinations can only be made by NCMUST*

Code of Federal Regulations (CFR)

- State Operations Manual Appendix PP revised 8/8/24:
<https://www.cms.gov/medicare/provider-enrollment-and-certification/guidanceforlawsandregulations/downloads/appendix-pp-state-operations-manual.pdf>
- Resident Assessment
 - Regulations F635-F646
- Comprehensive Resident Centered Care Plans
 - Regulations F655-F661

Regulation F641

Accuracy of Assessments

- ***The assessment must accurately reflect the resident's status.***
- ***INTENT:*** *To assure that each resident receives an accurate assessment, reflective of the resident's status at the time of the assessment, by staff qualified to assess relevant care areas and are knowledgeable about the resident's status, needs, strengths, and areas of decline.*

F641 Guidance

- Facilities are responsible for ensuring that all participants in the assessment process have the requisite knowledge to complete an accurate assessment.
- The assessment must represent an accurate picture of the resident's status during the observation period of the MDS... Be aware that different items on the MDS have different Observation Periods.
- When the MDS is completed, only those occurrences during the observation period will be captured on the assessment. In other words, if it did not occur during the observation period, it is not coded on the MDS.

Regulation F656

Develop/Implement Comprehensive Care Plan

- **The facility must develop and implement a comprehensive person-centered care plan for each resident, consistent with the resident that includes measurable objectives and timeframes to meet a resident's medical, nursing, and mental and psychosocial needs that are identified in the comprehensive assessment.**
- **The comprehensive care plan must describe the following —**
 - 1. The services that are to be furnished to attain or maintain the resident's highest practicable physical, mental, and psychosocial well-being and**
 - 2. Any services that would otherwise be required but are not provided due to the resident's exercise of rights, including the right to refuse treatment**
 - 3. Any specialized services or specialized rehabilitative services the nursing facility will provide as a result of PASARR recommendations.**

F656 Comprehensive Care Plan

- **4. In consultation with the resident and the resident's representative(s)—**
 - **A. The resident's goals for admission and desired outcomes.**
 - **B. The resident's preference and potential for future discharge. Facilities must document whether the resident's desire to return to the community was assessed and any referrals to local contact agencies and/or other appropriate entities, for this purpose.**
 - **C. Discharge plans in the comprehensive care plan, as appropriate, in accordance with the requirements set forth in paragraph (c) of this section.**
- **5. The services provided or arranged by the facility, as outlined by the comprehensive care plan, must be culturally-competent and trauma-informed.**

F656 Intent

- Each resident will have a person-centered comprehensive care plan developed and implemented to meet his or her preferences and goals, and address the resident's medical, physical, mental and psychosocial needs.

F656 Guidance

- Through the care planning process, facility staff must work with the resident and his/her representative, if applicable, to understand and meet the resident's preferences, choices and goals during their stay at the facility.
- The facility must establish, document and implement the care and services to be provided to each resident to assist in attaining or maintaining his or her highest practicable quality of life.
- Care planning drives the type of care and services that a resident receives.
- If care planning is not complete, or is inadequate, the consequences may negatively impact the resident's quality of life, as well as the quality of care and services received.

F657 Care Plan Timing and Revision

- *A comprehensive care plan must be—*
- *Developed within 7 days after completion of the comprehensive assessment/CAA.*
- *Prepared by an interdisciplinary team, that includes but is not limited to-- (A) The attending physician. (B) A registered nurse with responsibility for the resident. (C) A nurse aide with responsibility for the resident. (D) A member of food and nutrition services staff. (E) To the extent practicable, the participation of the resident and the resident's representative(s). An explanation must be included in a resident's medical record if the participation of the resident and their resident representative is determined not practicable for the development of the resident's care plan. (F) Other appropriate staff or professionals in disciplines as determined by the resident's needs or as requested by the resident.*
- *Reviewed and revised by the interdisciplinary team after each assessment, including both the comprehensive and quarterly review assessments.*

F657 Intent

- *To ensure the timeliness of each resident's person-centered, comprehensive care plan, and to ensure that the comprehensive care plan is reviewed and revised by an interdisciplinary team composed of individuals who have knowledge of the resident and his/her needs, and that each resident and resident representative, if applicable, is involved in developing the care plan and making decisions about his or her care.*

F657 Guidance

- GUIDANCE: Facility staff must develop the comprehensive care plan within seven days of the completion of the comprehensive assessment and review and revise the care plan after each assessment. “After each assessment” means after each assessment known as the Resident Assessment Instrument (RAI) or Minimum Data Set (MDS), except discharge assessments.
- For newly admitted residents, the comprehensive care plan must be completed within seven days of the completion of the comprehensive assessment and no more than 21 days after admission.

F644 Coordination of PASRR and Assessments

- **A facility must coordinate assessments with the pre-admission screening and resident review (PASARR) program under Medicaid to the maximum extent practicable to avoid duplicative testing and effort. Coordination includes:**
- **1. Incorporating the recommendations from the PASARR level II determination and the PASARR evaluation report into a resident's assessment, care planning, and transitions of care.**
- **2. Referring all level II residents and all residents with newly evident or possible serious mental disorder, intellectual disability, or a related condition for level II resident review upon a significant change in status assessment.**

F644 Intent

To ensure that the facility coordinates with the appropriate, State-designated authority, to ensure that individuals with a mental disorder, intellectual disability or a related condition receives care and services in the most integrated setting appropriate to their needs.

F644 Guidance

- The PASARR process requires that all applicants to Medicaid-certified nursing facilities be screened for possible serious mental disorders or intellectual disabilities and related conditions.
- This initial pre-screening is referred to as PASARR Level I and is completed prior to admission to a nursing facility.
- A negative Level I screen permits admission to proceed and ends the PASARR process unless a possible serious mental disorder or intellectual disability arises later.
- A positive Level I screen necessitates an in-depth evaluation of the individual by the state-designated authority, known as PASARR Level II, which must be conducted prior to admission to a nursing facility.

Validation Reports

- *Please* review your transmission validation reports regularly.
 - Reviewing will help you identify and correct errors
 - Reviewing will help prevent “missing assessments” and duplicate folders in the CMS data base
 - Reviewing will help ensure the facility will be paid

From RAI page 5-2

- When the transmission file is received by iQIES, the system performs a series of validation edits to evaluate whether or not the data submitted meet the required standards.
- MDS records are edited to verify that clinical responses are within valid ranges and are consistent, dates are reasonable, and records are in the proper order with regard to records that were previously accepted by iQIES for the same resident.
- The provider is notified of the results of this evaluation by error and warning messages on a Final Validation Report.
- All error and warning messages are detailed and explained in the Error Messages guide.

Section A:

Identification Information

Remember:

The CMS Database

matching process includes:

- First Name
- Last Name (A0500 D: Suffix: *Please use!*)
- Social Security Number
- Gender
- Date of Birth
- Please communicate regarding any changes to the resident's demographic information

Validation Report References

- **iQIES Resources are available at:**
<https://qtso.cms.gov/software/iqies/reference-manuals>
- **MDS 3.0 Provider User's Guide** is available at:
<https://www.cms.gov/medicare/quality-initiatives-patient-assessment-instruments/nursinghomequalityinits/mds30raimanual>
- **CASPER Reporting User's Guide For MDS Providers** is available at: <https://qtso.cms.gov/reference-and-manuals/casper-reporting-users-guide-mds-providers>

An interesting situation:

- A Short-Term (ST) resident had the same last name as an Established Resident (ER).
- Under ER's name, ST's entry tracker, admission, and discharge return not anticipated assessments were completed and transmitted. These assessments all had warnings on the final validation reports of change in first name, SSN, DOB, Medicare and Medicaid ID#s.
- When ER's next quarterly and OSA were completed and transmitted, there was a change in the resident ID number and a warning that the assessment was completed late due to no prior assessment found in the database.
- This warning and ID number change were noticed by the MDS nurse and along with the corporate nurse, couldn't figure out what happened to make the change.
- We were on the phone for 48 minutes figuring this out! Then I had to call iQIES to verify how to extract ST's assessments from ER's file.

Record 7	Status Accepted	Name [REDACTED]	XML File Name [REDACTED]
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Asmt_ID:	[REDACTED]	Name:	[REDACTED]
Res_Int_ID:	[REDACTED]	SSN:	[REDACTED]
A0200:	1	Medicare Num:	^
A0300A:	0	A0050:	NEW RECORD
A0310A:	99	Target Date:	07/23/2024
A0310C:	*	Attestation Date (X1100E):	^
A0310E:	0	Data Specs Version #:	3.01
A0310G:	^		
Item Subset Code:	NT		

MDS 3.0 Item(s):	A0500A, A0900
Item Values:	Old: [REDACTED] New: [REDACTED]
Message Number:	-1031
Message Type:	Warning
Message:	Resident Information Mismatch: Submitted value(s) for the item(s) listed do not match the values in the iQIES database. If the record was accepted, the resident information in the database was updated. Verify that the new information is correct.

MDS 3.0 Missing OBRA Assessment Report

Facility ID:

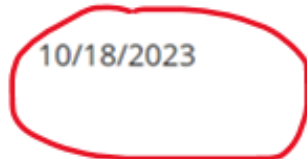
Facility Name:

City/State:



Report Run Date:

10/18/2023



Resident Identifiers:

Last Record Identifiers:

Resident
Internal ID

Resident
Name

SSN

Date of Birth

Gender

OBRA A0310A

PPS A0310B

Target Date

Filter

Filter

Filter

Filter

Filter

Filter

Filter

Filter



01

99

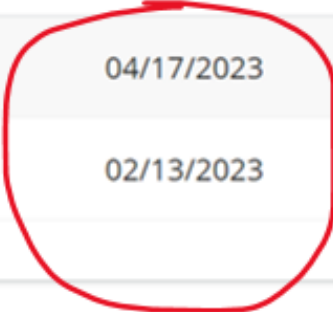
04/17/2023



99

99

02/13/2023



MDS 3.0 Missing OBRA Assessment Report

Facility ID: [Redacted]
 Facility Name: [Redacted]
 City/State: [Redacted]

Report Run Date: 08/13/2024

Resident Identifiers:					Last Record Identifiers:		
Resident Internal ID	Resident Name	SSN	Date of Birth	Gender	OBRA A0310A	PPS A0310B	Target Date
[Redacted]	[Redacted]	[Redacted]	[Redacted]	[Redacted]	99	01	05/30/2024

Patient Detail

Showing 4 of 4 assessments Merge/Split

<input type="checkbox"/>	Assessment ID	A0310A	A0310B	A0310F	A0310H	Assessment Type	Target Date	Submission ID	Submission Date
<input type="checkbox"/>	[Redacted]	99	99	01	0	Tracking	05/23/2024	[Redacted]	[Redacted]
<input type="checkbox"/>	[Redacted]	99	99	10	1	OBRA Discharge	05/29/2024	[Redacted]	[Redacted]
<input type="checkbox"/>	[Redacted]	99	01	99	0	PPS	05/30/2024	[Redacted]	[Redacted]
<input type="checkbox"/>	[Redacted]	-	-	-	-	Optional State Assessment	05/30/2024	[Redacted]	[Redacted]

MDS 3.0 Activity Report

Facility ID: [REDACTED]	Report Period: 10/01/2023 - 10/18/2023
Facility Name: [REDACTED]	Report Run Date: 10/18/2023
City/State: [REDACTED]	

Resident Intrnl ID/ SSN	Resident Name	Medicare Num	DOB/ Gender	A0310 A/B/C/D/F/G/H	ISC	Target Date	Subm Date	CALC MCR RUG
[REDACTED]	[REDACTED]	[REDACTED]	[REDACTED]	99/99/**/01/^/0	NT	09/25/2023	10/11/2023	*
[REDACTED]	[REDACTED]	[REDACTED]	[REDACTED]	02/99/**/99/^/0	NQ	09/15/2023	10/11/2023	KDSE

MDS 3.0 Missing OBRA Assessment Report

Displays the residents for whom the target date of the most recent OBRA assessment (other than a discharge or death record) is more than 138 days prior to the report run date. The report also includes residents for whom no OBRA record was submitted for a current episode that began more than 60 days prior to the report run date.

MDS 3.0 Activity Report

Displays a list of accepted assessments, tracking records and inactivation requests that were submitted by the requested facility(ies) for the time frame selected.

MDS 3.0 NH Final Validation Report

Displays detailed information regarding the records contained in the submission file for the facility. The report indicates whether the records were accepted or rejected and displays the warning and fatal errors for the records.



Report	Category	Last Run Date	Actions
MDS 3.0 NH Final Validation Report	Provider	08/07/2024 1:29 PM	Run Report
MDS 3.0 Missing OBRA Assessment Report	Provider	08/14/2024 11:01 AM	Run Report
MDS 3.0 Activity Report	Provider	06/07/2024 1:48 PM	Run Report
MDS 3.0 Roster Report	Provider	07/23/2024 2:49 PM	Run Report
SNF QRP Review and Correct Report	Quality Measure	08/19/2024 8:27 AM	Run Report
MDS 3.0 Resident-Level Quality Measure (QM) Report	Quality Measure	01/17/2024 11:21 AM	Run Report
SNF QRP Resident-Level Quality Measure (QM) Report	Quality Measure	08/17/2023 11:01 AM	Run Report
SNF QRP Provider Threshold Report	Quality Measure	08/17/2023 11:13 AM	Run Report

Choose “Spotlights and Announcements” for newest information for SNFs



- SNF Quality Reporting Program
- SNF Quality Reporting Program Spotlights & Announcements
- SNF Quality Reporting Program Measures and Technical Information
- SNF Quality Reporting Program Training

Skilled Nursing Facility (SNF) Quality Reporting Program (QRP) Public Reporting

Now available! Our new [Provider Data Catalog](#) makes it easier for you to search and download publicly reported data. We've also improved [Medicare's Compare sites](#).

Background

The Improving Medicare Post-Acute Care Transformation (IMPACT) Act of 2014 modified the Social Security Act requiring that SNFs be required to submit data for public reporting. In response, the Centers for Medicare & Medicaid Services (CMS) established the SNF QRP and authorized the

Skilled Nursing Facility (SNF) Quality Reporting Program (QRP) Measures and Technical Information

- <https://www.cms.gov/medicare/quality/snf-quality-reporting-program/measures-and-technical-information>

CMS Training Videos

- <https://www.cms.gov/Medicare/Quality-Initiatives-Patient-Assessment-Instruments/NursingHomeQualityInits/Skilled-Nursing-Facility-Quality-Reporting-Program/SNF-Quality-Reporting-Program-Training.html>
- <https://www.cms.gov/medicare/quality-initiatives-patient-assessment-instruments/nursinghomequalityinits/skilled-nursing-facility-quality-reporting-program/snf-quality-reporting-program-spotlights-and-announcements>
- <https://www.youtube.com/playlist?list=PLaV7m2-zFKphoXW6cc3NwUfxra0A1LYDi>
- <https://www.cms.gov/medicare/quality/snf-quality-reporting-program/training>
- Accessed 9/20/24

Another Helpful Site

CMS Nursing Home Resource Center

<https://www.cms.gov/nursing-homes>

Helpful Resource for Documentation

- Medicare Benefit Policy Chapter 8 Coverage of SNF Services:
- <https://www.cms.gov/Regulations-and-Guidance/Guidance/Manuals/Downloads/bp102c08pdf.pdf>
- NC Medicaid, Nursing Facility Services Clinical Coverage Policy:
- <https://Medicaid.ncdhhs.gov/media/12254/open>
- Myers and Stauffer:
- <https://myersandstauffer.com/client-portal/north-carolina/>

MDS RAI Manual Version 1.19.1 effective October 2024

- MDS RAI Manual version 1.19.1 and Item Sets available: <https://www.cms.gov/medicare/quality/nursing-home-improvement/resident-assessment-instrument-manual>
- Final Rule: <https://www.federalregister.gov/documents/2023/08/07/2023-16249/medicare-program-prospective-payment-system-and-consolidated-billing-for-skilled-nursing-facilities>

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Thank you!

- Thank you for all the work you do to ensure the care, comfort and safety of our most vulnerable in society. This is not an easy job you do, and it must come from the heart. Weariness and frustration can easily become your best friends, but don't let them take over! Know that you are not alone in your work. Reach out, make friends and contacts who will encourage your soul. Please know that you are welcome to call or email me anytime.
Sincerely, Janet