

Overview of IPFQR Program Resources, Part 2

Evette Robinson, MPH, CPHQ

Program Lead

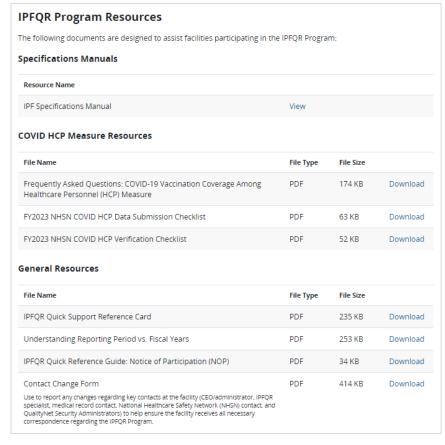
Inpatient Psychiatric Facility Quality Reporting (IPFQR) Program Inpatient Value, Incentives, and Quality Reporting (VIQR)

Outreach and Education Support Contractor

June, 2022

Introduction

Part 1 of this webinar series included a review of the resources available for download from the QualityNet IPFQR Program Resources web page.



IPFQR Program Questions and Answers Tool	View		
Click on the link to search for posted question and answer pairs as well as submit any new	view		
questions that are not already addressed in the Q&A tool or in a published summary of question and answers.			
iscal Year (FY) 2023 Payment Determination			
File Name	File Type	File Size	
FY 2023 IPFQR Program Guide	PDF	325 KB	Download
IPFQR Program Measures for FY 2023	PDF	109 KB	Download
IPFQR Quick Reference Guide: FY 2023 DACA	PDF	49 KB	Download
iscal Year (FY) 2024 Payment Determination			
File Name	File Type	File Size	
FY 2024 IPFQR Program Guide	PDF	325 KB	Download
IPFQR Program Measures for FY 2024	PDF	111 KB	Download
IPFQR Quick Reference Guide: FY 2024 DACA	PDF	103 KB	Download
ata Submission and Verification Checklists			

Purpose

This presentation will help participants navigate the QualityNet website to locate the Specifications Manual for National Inpatient Psychiatric Facility Quality Measures (IPF Specifications Manual) and answer several commonly asked questions related to the manual.

Objectives

At the conclusion of this presentation, attendees will be able to locate the IPF Specifications Manual on QualityNet and leverage the contents of the manual to optimize success in the IPFQR Program.

Webinar Questions

Please email any questions that are pertinent to the webinar topic to WebinarQuestions@hsag.com.

- Write "IPF Program Resources, Part 2" in the subject line.
- If your question pertains to a specific slide, include the slide number in the body of the email.

Overview of IPFQR Program Resources, Part 2

Navigating to the IPF Specifications Manual Web Page

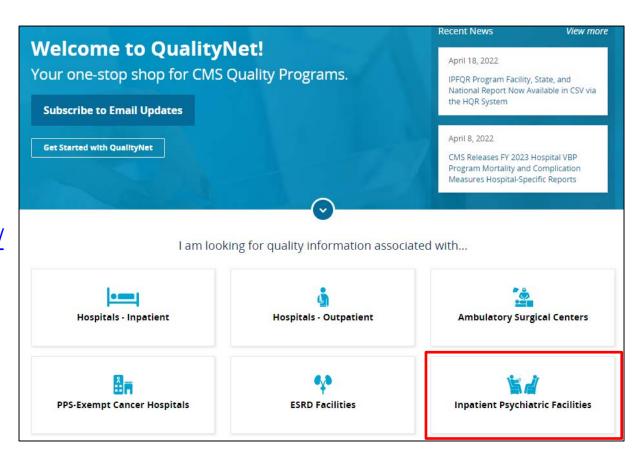
QualityNet Home Page

There are a few ways to access the IPF Specifications Manual.

The simplest way is via the direct URL for the IPF Specifications Manuals web page:

https://qualitynet.cms.gov/ ipf/specificationsmanuals#tab1

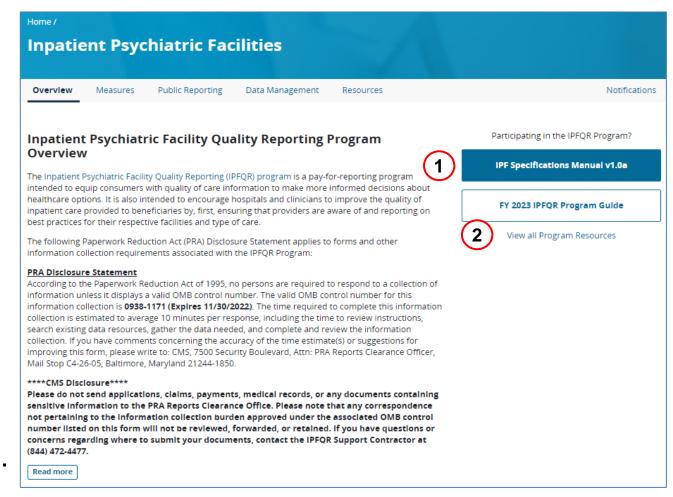
To navigate to the web page from the QualityNet home page, click on the icon for Inpatient Psychiatric Facilities.



Inpatient Psychiatric Facilities Overview Web Page

From the IPF
Overview web page
you can access the
manual by clicking
the blue button (1),
IPF Specifications
Manual. This will
take you to the
latest version.

As shown in Part 1, you can also access the manual (2) from the IPFQR Program Resources web page.



Inpatient Psychiatric Facilities Overview Web Page

At the bottom of the IPF Overview web page is a Learn more button (3).



Inpatient Psychiatric Facility Quality Reporting Program Overview Web Page

On the <u>IPFQR Overview</u> web page, where you can access the latest version of the IPF Specifications Manual under Key Documents on the right side of the web page.

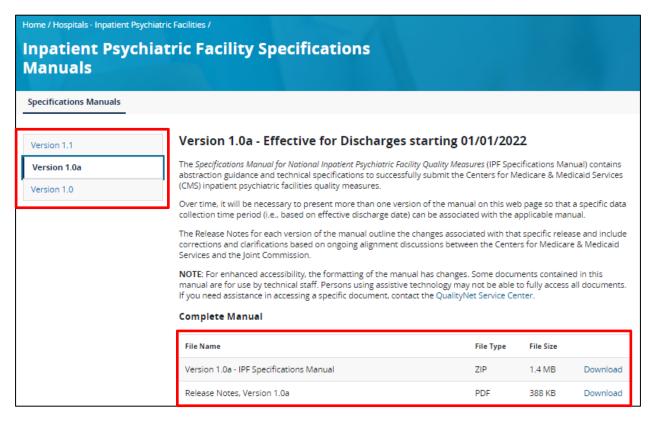


Overview of IPFQR Program Resources, Part 2

IPF Specifications Manual

IPF Specifications Manual Web Page

By default, the most recently published version of the manual is displayed. You can access prior versions from the menu on the left side of the screen. This presentation focuses on the IPF Specification Manual Version 1.0a.



IPF Specifications Manual

When you download and open the ZIP folder, you will see three files:

- 1. IPF Specifications Manual
- 2. IPF Clinical Data XML File Layout
- 3. IPF Non-Measure XML File Layout
 - IPF Specifications Manual v1.0a_FINAL(508)
 - IPF_ClinicalDataXMLFileLayout_v1.0a_Final
 - IPF_Non-MeasureXMLFileLayout_v1.0a_Final

IPF Specifications Manual

Specifications Manual for Inpatient Psychiatric Facility Quality Measures

Version 1.0a Effective date: January 1, 2022 discharges (All data that are to be reported to CMS in calendar year 2023)

Table of Contents	
Introduction	
Specifications Manual for National Inpatient Psychiatric Facility Quality Measures	
Section 1: Data Dictionary	
Introduction to the Data Dictionary	
Alphabetical Data Dictionary	
Section 2: Measure Information	
Measures Stewarded by The Joint Commission	
Screening for Metabolic Disorders Measure	
SMD Data Element List	
Screening for Metabolic Disorders (SMD) Initial Patient Population	
Screening for Metabolic Disorders (SMD) Sample Size Requirements	7
Measure Information Form (MIF) and Flowchart (Algorithm) SMD78	8
Transition Record Measures	4
TR Data Element List	4
Transition Record (TR) Measures' Initial Patient Population	6
Transition Record (TR) Measures' Sample Size Requirements	6
Measure Information Form (MIF) and Flowchart (Algorithm) TR	7
Section 3: Missing and Invalid Data	0
Section 4: Population and Sampling Specifications	3
Section 5: Data Transmission	6
Introduction	6
CMS Data Transmission	6
Guidelines for Submission of Data	6
Transmission Data Processing Flow11	4
Outcome Measures (Claims Based)114	4
Appendices 11	7
Appendix A: ICD-10 Code Tables11	7
Appendix B: Medication Tables	8
Appendix C: Crosswalk Tables for Non-Measure Data	9
Appendix D: Glossary of Terms	6
Appendix E: Overview of Measure Information Form and Flowchart Formats	6
Appendix F: Resources	2
Appendix G: Psychiatric Advance Directives (PAD)	4
Appendix H: Measure Name Crosswalk	
Appendix I: Preview Section	7
••	

IPF Specifications Manual

		Inpatient Psychiatric Facility (IPF) Clinica	ıl Data XML i	File Layout						IPF	= C	lir	nic	al Data XML	
Note: This file is to be u	sed as posted Marc	h 18, 2022, and CMS is not responsible for pote	ntial errors and	d issues arising from	n modification	ns made by exte	ernal partie	PS.						The state of the s	
XML Element	Attributes	Description	Data Element	Valid Values	Data Type	Field Size	Data Require (CMS	ed		File	e L	.ay	yo u	ıt: <u>Elements</u>	
A header is OPTIONA xml version="1.0" end</td <td></td> <td>of each XML file as follows:</td> <td></td> <td></td> <td></td> <td></td> <td>ı</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>Int</td> <td>F Clinical Data XML File Layout</td> <td></td>		of each XML file as follows:					ı						Int	F Clinical Data XML File Layout	
<submission></submission>	Opening tag is requ	ired.												•	
	type	Describes the setting for which data is being		IPF	Character		Yes	0						MS is not responsible for potential errors and issues arising from modifications made by external parties.	Analisable
	data version	Describes the type of data being submitted. The version of the file layout.	N/A N/A	1.0	Character Character		Yes Yes	Question	Question/ Field Name	Data Type	Size	occui	r Answe		Applicable Measure(s)
	action-code	Describes the intended action of the file being			Character		Yes	including physician	247CONTACT	Alphanumeric	1	1	Υ		TR1
<file-audit-data></file-audit-data>	Opening tag for	Note: This tag and the entire <file-audit-data></file-audit-data>	section are	optional in the XML	document.	If submitted, t	his tag	for emergencies related to inpatient							
Sub-element of the	file audit data	contains no data.						stay					N	No	
submission data element		Required if sub-elements are included.													
<create-date></create-date>	Each element must have a closing tag that is the same as the opening tag but with a forward slash.								Suggested Data Collection Question: Does the transition record contain documentation that the patient has an advance care plan or a reason for not providing an advan						
Sub-element of the file	Example with da							or surrogate decision maker	ADPADSDM	Alphanumeric	1	1	ly .	Yes	TR1
audit data element		10-2021	laura.	luu pp vooor	In a	T ₄ 0		decision maker documented OR	ADI ADODIII	Aprilandino	·	l '	Ι΄.		
	None	The month, day, and year the XML file was created	N/A	MM-DD-YYYY Must be a valid	Date	10		documented reason					N	No	
		Created		date:				for not providing					"	NO	
				MM (01-12)				advance care plan							
				DD (01-31) YYYY (20xx)				Alcohol Use Status	Suggested Data Collect			patient	's alcohol i		
<create-time></create-time>	Each element must	have a closing tag that is the same as the open	ning tag but w		sh.	1	-		ALCSTATUS	Alphanumeric	1	1	1	The patient is screened with a validated tool within the first day of admission (by end of Day A 1) and the score on the alcohol screen indicates no or low risk of alcohol related problems.	All SUB Measures
Sub-element of the file	Example with data:						ľ						2	The patient was screened with a validated tool within the first day of admission (by end of	
audit data element	<create-time>23</create-time>					1_								Day 1) and the score on the alcohol screen indicates unhealthy alcohol use (moderate or	
	None	The hour and minutes representing the time the file was created.	N/A	HH:MM (Military format with or	Time	5	No						3	high risk) benefitting from brief intervention. The patient was screened with a non-validated tool within the first day of admission (by end	
		the life was created.		without colon)									-	of Day 1) and the score on the alcohol screen indicates no or low risk of alcohol related	
				HH (00-23)									4	problems. The patient was screened with a non-validated tool within the first day of admission (by end	
<create-by></create-by>	Each element must	have a closing tag that is the same as the oper	ning tag but w	MM (00-59)	sh		\vdash						*	of Day 1) and the score on the alcohol screen indicates unhealthy alcohol use (moderate or	
Sub-element of the file	Example with da	a:	ing tag but t	This distributed on the									<u> </u>	high risk) benefitting from brief intervention.	
audit data element	<create-by>Vend</create-by>		I	I		1							5	The patient refused the screen for alcohol use within the first day of admission (by end of Day 1).	
	None	The entity who created the file	N/A	Up to 50 letters, numbers, and/or	Character	50	No						6	The patient was not screened for alcohol use within the first day of admission (by end of	
				special									7	Day 1) or unable to determine from medical record documentation. The patient was not screened for alcohol use within the first day of admission (by end of	
				characters can									1	Day 1) because of cognitive impairment.	
				be entered. Only the following				Appropriate		•			*		
				special				Justification for		tion Question: Is t	here doc	umenta	ation in the	e medical record of appropriate justification for the patient being discharged on two or more antipsychoti	
				characters will be allowed:				Multiple	JSTPSYCHMD	Alphanumeric	1	1	1		HBIPS-5
				~ ! @ # \$ %				Antipsychotic Medications					2	multiple trials of monotherapy. 2 The medical record contains documentation of a recommended plan to taper to	
				^ * () _ + { }				modications						monotherapy due to previous use of multiple antipsychotic medications OR documentation of	
				[] : ? ` - = []									2	a cross-taper in progress at the time of discharge. 3 The medical record contains documentation of augmentation of Clozapine.	
				\; '., / and									4	4 The medical record contains documentation of a justification other than those listed in	
<pre><version> Sub-element of the file</version></pre>	Each element must Example with da	have a closing tag that is the same as the oper	ning tag but w	vith a forward slas	sh.									Allowable Values 1-3.	
audit data element	<pre><version>1.0</version></pre>												5	5 The medical record does not contain documentation supporting the reason for being discharged on two or more antipsychotic medications OR unable to determine from medical	
	None	The version of the file being submitted	N/A	Up to 20 letters,	Character	20	No							record documentation.	
				numbers, and/or	1			Blood Glucose	Suggested Data Collect	tion Question: Is t	here doc	uments	ation of a n	numerical value of blood glucose in the patient's medical record during this stay or at any time during the	12 months
				special characters can				Dioda Ciacco	prior	non adoction. Io				numerical radio of blood glacoboo in the patient of incurcal record during the stay of at any time during the	TE IIIOIIIIIO
				ho ontored Only					BLDGLUCOSE	Alphanumeric	1	1	Υ	Yes	SMD
← → Elem	ents Detail	Elements Info 🕒											N	No	
					Blood Pressure	Suggested Data Collect time during the 12 mont			blood	pressure	(numerical systolic and diastolic values in mmHg) documented in the patient's medical record during this	stay or at any			
	וחו	F Clinical Data	o VI	AL C::		1	ŀ		BLDPRESS	Alphanumeric	1	1	Υ	Yes	SMD
	IPI	- Cimicai Dati	a All	∕IL FII	e								N	No	
	100							< → E	lements Detail E	lements Info	((+)		1	
	Lay	out: <u>Detail Ele</u>	<u>eme</u>	nts Ir	110										

How will I know when a new version of the manual is available for download?

Are separate XML files required for each episode of care?

What changed from the previous version to the current version of the manual?

Where can I find a complete list of the 11 elements that must be in the TR-1 measure?

Where is information about the COVID-19 Vaccination Coverage Among HCP measure?

How do IPFs abstract for the advance care plan data element for minors?

Do the following examples meet the Reason for IPF Admission data element?

Can parts of the transition record be electronic and the rest on paper?

Can I upload a ZIP file containing 15k XML files into the HQR System?

Why isn't there an IPFQR Program Manual after v7.0a?

Due to CMS' decision to transition from aggregate to patient-level reporting of measure data for the IPFQR Program (per the FY 2022 IPF PPS Final Rule: https://www.govinfo.gov/content/pkg/FR-2021-08-04/pdf/2021-16336.pdf), CMS decided to discontinue producing the IPFQR Program Manual. Instead, CMS created the IPF Specifications Manual and several supporting resources, which will be used going forward. These documents are modeled after those used by the Hospital IQR Program.

How will I know when a new version of the manual is available for download?

CMS communicates the availability of the IPF Specifications Manual via email notifications distributed to IPFQR Program Listserve subscribers. We recommend that you sign up for the IPFQR Program Listserve (https://qualitynet.cms.gov /listservsignup), if you have not already, to ensure that you receive timely communications regarding the availability of the manual, other IPFQR Program resources, and updates.

What changed from the previous version to the current version of the IPF Specifications Manual?

A Release Notes document is posted each time a new version of the manual is published. Where possible, the document includes the page number where information was added or changed. The document also states when information was removed. You can download the Release Notes from the QualityNet Inpatient Psychiatric Facility Specifications
Manuals web page: https://qualitynet.cms.gov/ipf/specifications-manuals

Why is the Timely Transmission of Transition Record (TR-2) measure in the manual for CY 2022 discharges if the measure was discontinued after CY 2021?

As stated in footnote 2 on page 87 of the IPF Specifications Manual, v1.0a:

"Eligible IPFs will collect Timely Transmission of Transition Record (TR-2) measure data through December 31, 2021 and report the data to CMS for the IPFQR Program for the last time during the summer 2022 data submission period. This IPF Specifications Manual provides guidance for CY 2022 discharges; however, this measure is included because it is the only resource available to providers who choose to voluntarily report TR-2 measure data at the patient-level during the summer of 2022."

How do I abstract for the TR-1 and TR-2 measures if the patient has multiple admissions to the IPF and the visits are combined into one encounter?

A transition record must be created for each discharge from the IPF and abstracted accordingly, not based on how the patient encounters are billed. Refer to the Episode of Care section on page 7 of the IPF Specifications Manual, v1.0a:

"For the Transition Record measures, abstract each discharge from the IPF separately, regardless of whether the patient was discharged from the IPF to home, to another unit within the same facility, or to a different inpatient facility. If a patient is transferred from an IPF unit to another IPF unit within the same healthcare system and the IPF units share the same CCN, this should be abstracted as one episode of care."

☐ KB0017308: Multiple admissions/discharges

Where can I find a complete list of the 11 data elements that must be in the transition record?

Th	e 11 required data elements of the transition record are:
	24-hour/7-day contact information including physician for emergencies related to inpatient stay
	Advance directives or surrogate decision maker documented OR documented reason for not providing advance care plan
	Contact information for obtaining results of studies pending at discharge
	Current medication list
	Major procedures and tests performed during inpatient stay and summary of results
	Patient instructions
	Plan for follow-up care
	Primary physician, other health care professional, or site designated for follow-up care
	Principal diagnosis at discharge
	Reason for inpatient admission

6/2022

Studies pending at discharge (or documentation that no studies are pending)

Our after-visit summary (AVS) and discharge summary together include all 11 data elements.

Can the transition record be multiple documents?

The transition record must be clearly identifiable as a single document containing all 11 data elements and it can be multiple pages long. If one or more elements is missing from the transition record - even though the information may be available elsewhere in the patient's medical record (e.g., via the patient portal) or provided to the patient as a separate document - the transition record is considered incomplete, and the case will not be included in the numerator for the Transition Record measures.

- □ KB0017254: Transition record 11 elements on different forms
- □ KB0017282: All 11 elements must be in the transition record
- ☐ <u>KB0017218</u>: Transition Record Missing element

Must the contact person for the 24-hour/7-day contact information including physician for emergencies related to inpatient stay data element be a physician?

As stated in the definition of the 24-hour/7-day contact information, including physician for emergencies related to inpatient stay data element on pages 16 and 146 of the Specifications Manual for National Inpatient Psychiatric Facility Quality Measures, v1.0a, it is appropriate to abstract "Y (Yes)" when the transition record includes documentation regarding the "Physician, health care team member, or other health care personnel who has access to medical records and other information concerning the inpatient stay and who could be contacted regarding emergencies related to the stay." (https://qualitynet.cms.gov/ipf/specifications-manuals).

Our hospital uses a signature page to indicate the transition record was discussed with and provided to the patient and/or caregiver.

What should we abstract if they refuse to sign?

Signatures are not required and do not indicate in and of themselves whether the transition record was discussed with and provided to the patient, the patient's caregiver, or both. If there is documentation anywhere in the medical record indicating that the transition record was discussed with and provided to the patient or the patient's caregiver at discharge, then it is acceptable to abstract value "1 Transition record was discussed with and provided to the patient and/or caregiver at discharge" for the *Transition Record Discussed and Provided* data element, as described on page 71 of the IPF Specifications Manual, v1.0a.

If there is no such documentation in the medical record and the patient or the patient's caregiver did not sign the signature page, then abstract value "3 Transition record was not discussed with and/or provided to the patient and/or caregiver at discharge or Unable to Determine (UTD) from the medical record documentation" for the *Transition Record Discussed and Provided* data element.

☐ KB0017249: Documentation that transition record was discussed

Does the transmission of all 11 elements in the transition record to the inpatient facility satisfy the measure, or does it have to be verbally discussed with the receiving facility?

If a patient is discharged from an IPF to an inpatient facility, then the *Four Elements Discussed with Receiving Inpatient Facility* data element must be met.

Documentation of verbal communication regarding the four elements (at a minimum), is required to abstract "Y (Yes)", as defined for the *Four Elements Discussed with Receiving Inpatient Facility* data element on page 38 of the IPF Specifications Manual, v1.0a. The discussion can occur during verbal report when the patient transitions to a medical floor (for example) or by phone to the receiving inpatient facility.

- □ KB0017169: Transition record for patient transferred to medical floor
- ☐ KB0017170: Phone conversation between discharging and receiving inpatient facilities
- ☐ KB0017313: Documentation of four elements discussed with receiving facility
- ☐ KB0017283: Elements to be discussed with receiving inpatient facility

When tests were completed during the visit, do the results need to be documented in the transition record, or is a blanket statement that the results were discussed with the patient and/or caregiver sufficient?

No, a blanket statement is not sufficient. Documentation of the specific noteworthy test(s) performed, and the results in the transition record, is required to abstract "Y (Yes)" for the *Major Procedures and Tests Performed During Inpatient Stay and Summary of Results* data element, unless there is documentation that none were performed during the IPF stay. Refer to the second bullet in the Notes for Abstraction for this element on page 44 of the IPF Specifications Manual, v1.0a:

"The name and results of noteworthy procedures and tests performed during the IPF stay must be documented to meet this element, if applicable."

Also, note the following abstraction guidance on page 9 of the IPF specifications manual:

"The medical record must be abstracted as documented (taken at "face value"). Information should not be added after the fact and assumptions should not be made to meet a measure. Documentation is not to be added at the time of abstraction to ensure the passing of measures for the IPFQR Program. When the value documented is obviously in error (not a valid format/range or outside of the parameters for the data element) and no other documentation is found that provides this information, the abstractor should select 'UTD.'"

If the patient's active problem list is documented in the transition record, is this sufficient to abstract "Y (Yes)" for the *Principal Diagnosis at Discharge* data element?

No. Refer to the second bullet in the Notes for Abstraction for the *Principal Diagnosis at Discharge* data element on page 63 of the IPF Specifications Manual, v1.0a, which states:

"The principal diagnosis must be clearly identified in the transition record to meet this data element. A problem list cannot be used to meet this element."

If the discharge diagnosis list is numbered but there is no distinguishing principal diagnosis, will this meet the *Principal Diagnosis at Discharge* data element?

No. The second bullet in the Notes for Abstraction for the *Principal Diagnosis at Discharge* data element on page 62 of the IPF Specifications Manual, v1.0a, states:

"The principal diagnosis must be clearly identified in the transition record to meet this data element."

Listing more than one diagnosis under "Principal Diagnosis" in the transition record introduces ambiguity as to which one is the final principal diagnosis and thereby would not meet the *Principal Diagnosis at Discharge* data element. If the transition record clearly labels the final principal diagnosis at discharge and lists only one diagnosis under that label (regardless of other diagnoses listed elsewhere in the transition record), then the *Principal diagnosis at Discharge* data element is met. If, when abstracting, there is ambiguity or doubt as to whether a data element is met, then abstract "No" for the data element.

What types of facilities meet value "1 Home" for the *IPF Discharge Disposition* data element?

If the patient was discharged to a place where the patient will no longer receive inpatient-level healthcare services (such as a home residence or homeless shelter), then it is appropriate to abstract Value "1 Home" for the *IPF Discharge Disposition* data element.

What types of facilities meet value "2 Inpatient Facility" for the *IPF Discharge Disposition* data element?

To determine whether the patient was discharged to an inpatient facility, refer to the second bullet in the Notes for Abstraction for the *IPF Discharge Disposition* data element on page 40 of the IPF Specifications Manual, v1.0a, which states: "If the patient is discharged to a location where the patient will receive inpatient-level health care services, then abstract value '2 Inpatient Facility'."

A few examples (though not inclusive) are included in the "inpatient facility" definition on page 150 of the manual: "hospital inpatient or observation, skilled nursing facility (SNF), rehabilitation facility, or IPF."

6/2022 31

To abstract value 3 for the *IPF Discharge Disposition* data element, how can I tell if a patient discontinued care?

It is appropriate to abstract value 3 for the *IPF Discharge Disposition* data element when there is documentation in the medical record that the patient died, left against medical advice, or discontinued care. Per the definition of the term "discontinued care" on page 149 of the IPF Specifications Manual, v1.0a, it "includes elopement and failure to return from leave", both of which are described in the definition.

□ KB0017168: Definition of discontinued care

Advance directives are not enforceable in my state.

Does documentation of the law meet the data element?

No, because the psychiatric and non-psychiatric (medical) advance directives or the designation of a surrogate decision maker can still be completed even if it is not enforceable.

- □ <u>KB0017236</u>: Psychiatric Advance Directive (PAD) not recognized in some states
- □ KB0017315: Advance care plan for patients under the age of 18

Does the surrogate decision maker need to be identified by first and last name to meet the data element?

There are no format requirements for the name of the surrogate decision maker. Per the last sentence in the definition of the term "Surrogate decision maker" on page 154 of the IPF Specifications Manual, v1.0a:

"The surrogate decision maker must be identified in the transition record by name and telephone number."

Documentation of a descriptor and first name (e.g., "daughter Sue") and the surrogate decision maker's contact number would meet this requirement. Ideally, the surrogate decision maker would be identified by first and last name in the transition record to reduce/avoid confusion, particularly if the patient and the surrogate decision maker share the same or similar names, but the inclusion of the telephone number is also important to ensure that the person can be contacted.

What do I need to document to meet the Reason for IPF Admission data element?

To abstract "Y (Yes)" for the *Reason for IPF Admission* data element, the transition record must include documentation that states:

- How the patient came to be admitted to the IPF (e.g., self-admit, brought in by family member, transported by police, admitted through the emergency department) OR
- Why the patient was admitted to the IPF (e.g., behaviors and symptoms that led to IPF admission) AND
- The triggering or precipitating event that led to IPF admission, if applicable.

A triggering event should be included if there was one. In the absence of a triggering event, a clear description of how and/or why the patient was admitted is sufficient.

Can you provide a specific example of documentation for the Reason for IPF Admission data element? Is "danger to self" sufficient to abstract "Yes"?

Specific examples are provided in the Notes for Abstraction for the *Reason for IPF Admission* data element on page 64 of the IPF Specifications Manual, v1.0a. The sample documentation demonstrates how an IPF can succinctly include all the information that is most useful for this data element. Documentation of "danger to self" is too vague, but it can be conveyed with more precision, as shown in the following example from page 64 of the manual:

"Jane Doe was admitted with a 2-month history of an increasingly depressed mood, difficulty sleeping and suicidal thoughts with a plan to take an overdose. Recent events include poor adherence with antidepressant treatment, becoming homeless and conflict with family that led them to contact police."

I reviewed the specifications and the Notes for Abstraction for the *Reason for IPF Admission* data element. Can you help me understand how to determine if the documentation is sufficient?

Here is a breakdown of how the first example provided on page 64 of the IPF Specifications Manual, v1.0a, meets the *Reason for IPF Admission* data element.

- Why: "increasingly depressed mood, difficulty sleeping and suicidal thoughts with a plan to take an overdose"
- How: "police"

 Triggering/Precipitating Event(s): "poor adherence with antidepressant treatment, becoming homeless and conflict with family"

Would documentation of "Needs detox; psychosis or thought disorder" meet the *Reason for IPF Admission* data element?

No, the documentation provided does not meet the *Reason for IPF Admission* data element. The documentation is missing a clear description of how and/or why the patient was admitted to the IPF. Documentation of "Psychosis or thought disorder" is not sufficient per the last sentence in the definition of the data element: "A diagnosis or a list of symptoms alone is not sufficient." Also, documentation of "needs detox" is too vague.

Additional detail around how the patient came to the IPF (e.g., self-admit, brought in by family, etc.), the behaviors that led to admission (e.g., patient expressed auditory and visual hallucinations to family member), and a description of the triggering event (e.g., patient attempted detox from heroin) would meet the data element.

Are separate XML files required for each episode of care?

Yes. Refer to the guidance on page 107 in the IPF Specifications Manual, v1.0a: "Each case must have a separate XML file." Exceptions pertaining to the Transition Record and HBIPS event measures are also described.

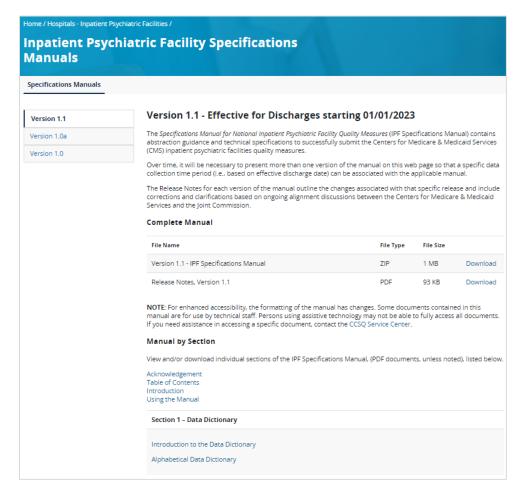
- Transition Record Measures: Each discharge from the IPF must be abstracted as a separate episode of care, not based on how the inpatient stay was billed; regardless of whether the patient was discharged from the IPF to home, to another unit within the same facility, or to a different inpatient facility.
- HBIPS-2 and HBIPS-3: If a patient has multiple events specific patient identifiers must match for each event record transmitted.

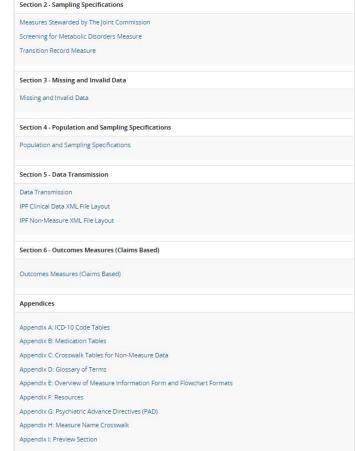
Can a ZIP folder containing 15k XML files be uploaded into the HQR System?

Yes, a single ZIP folder containing 15,000 XML files for the reporting period will be acceptable.

When will the CY 2023 specifications be available?

Version 1.1 of the IPF Specifications Manual was posted to the QualityNet website on May 27, 2022.





6/2022 41

Overview of IPFQR Program Resources, Part 2

What's next?

Future Webinar Topics

Overview of IPFQR Program Resources, Part 3 (On Demand)

Specifications Manual for Inpatient Psychiatric Facility

Quality Measures, Version 1.1

Keys to Successful FY 2023 Data Reporting



Future webinar titles, dates, and times will be communicated via the IPFQR Program ListServe.

Overview of IPFQR Program Resources, Part 2

Helpful Resources

Acronyms

AVS	after visit summary	IPF	inpatient psychiatric facility
CCN	CMS Certification Number	IPFQR	Inpatient Psychiatric Facility Quality Reporting
CDC	Centers for Disease Control and Prevention	IQR	Inpatient Quality Reporting
CMS	Centers for Medicare & Medicaid Services	KB	knowledge base
CY	calendar year	PAD	psychiatric advance directive
EVT	event	PPS	prospective payment system
FY	fiscal year	Q	quarter
HBIPS	Hospital-based Inpatient Psychiatric Services	SNF	skilled nursing facility
HBIPS-2	Hours of Physical Restraint Use	TR-1	Transition Record with Specified Elements Received by Discharged Patients
HBIPS-3	Hours of Seclusion Use	TR-2	Timely Transmission of Transition Record
HCP	healthcare personnel	UTD	unable to determine
HQR	Hospital Quality Reporting	VIQR	Value, Incentives, and Quality Reporting

Helpful Resources



Helpful Resources

Stay up to date...



...and get answers to your questions.









Webinar Questions

Please email any questions that are pertinent to the webinar topic to WebinarQuestions@hsag.com.

- Write "IPF Program Resources, Part 2" in the subject line.
- If your question pertains to a specific slide, include the slide number in the body of the email.

Overview of IPFQR Program Resources, Part 2

Thank You!

Disclaimer

This presentation was current at the time of publication and/or upload onto the Quality Reporting Center and QualityNet websites. Medicare policy changes frequently. Any links to Medicare online source documents are for reference use only. In the case that Medicare policy, requirements, or guidance related to this presentation change following the date of posting, this presentation will not necessarily reflect those changes; given that it will remain as an archived copy, it will not be updated.

This presentation was prepared as a service to the public and is not intended to grant rights or impose obligations. Any references or links to statutes, regulations, and/or other policy materials included in the presentation are provided as summary information. No material contained therein is intended to take the place of either written laws or regulations. In the event of any conflict between the information provided by the presentation and any information included in any Medicare rules and/or regulations, the rules and regulations shall govern. The specific statutes, regulations, and other interpretive materials should be reviewed independently for a full and accurate statement of their contents.