Radiation Treatment QBP Operational Report Dashboard on HSI

External User Training

Ontario Health – Funding Unit

Ontario Health

Introducing the Radiation Treatment (RT) Dashboard

- Consolidates all information and data available on Radiation Treatment activity and funding including:
  - Bundled services methodology
  - Patient care activities
  - Infrastructure requirements for operating RT programs

- The RT dashboard will be launched on Ontario Health's Health System Insights (HSI) platform late 2026/early 2027
- The dashboard will include historical data, dating back to 2022-2023 FY



Benefits of Switching RT Reports to HSI

- Most up-to-date data available
- Readily access historical data
- Secure portal to ensure no breaches
- Direct access for facilities to track reporting
- Centralized platform to reduce errors in reporting

Training Objectives

By the end of this session, you will be able to:

1. Understand how to access the dashboard
2. Navigate through different years of RT reports
3. Understand how to filter through data tabs
4. Know how to get support when you need it

Platform Access

- Registration Process
- Logging into the HSI Platform

Gain Access to the HSI Platform

Set-up ONE® ID Account

- Contact your facility ONE® ID Local Registration Authority (LRA) to request enrollment to the HSI Service

Log in to HSI

- Navigate to https://hsi.ontariohealth.ca
- Click ONE® ID Button and log in with ONE® ID credentials
- Complete the HSI Welcome Wizard
  - Acknowledge the End Sure License Agreement
  - Complete profile details and select dashboards

Contact your HSI FRA

- New and existing HSI Users must contact the facility HSI Facility Role Administrator (HSI FRA) to approve access to requested dashboards

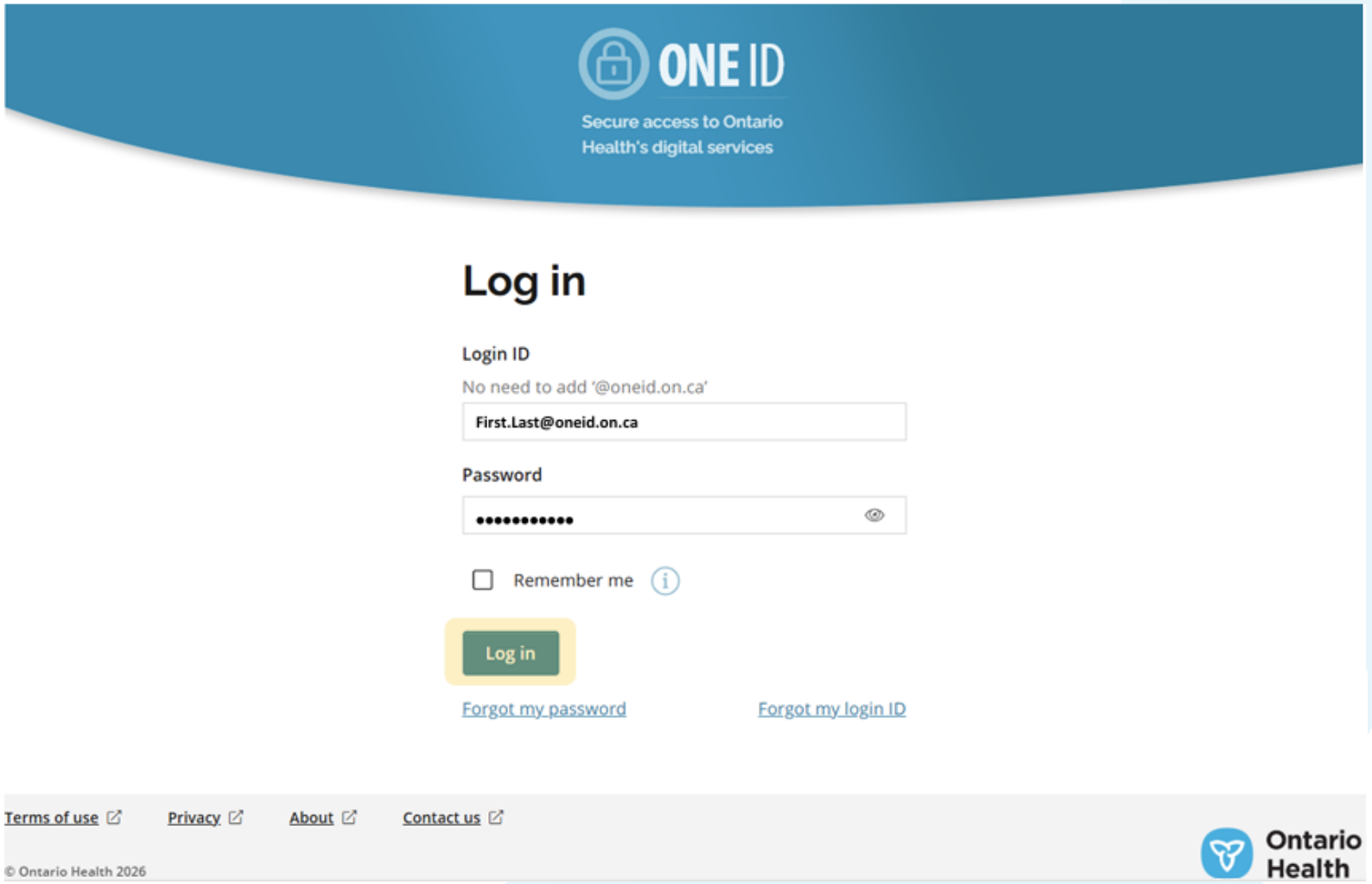
Note: For contact information about your facility's ONE® ID LRA or HSI FRA, please contact: itservicedesk@ontariohealth.ca

Accessing the HSI Platform

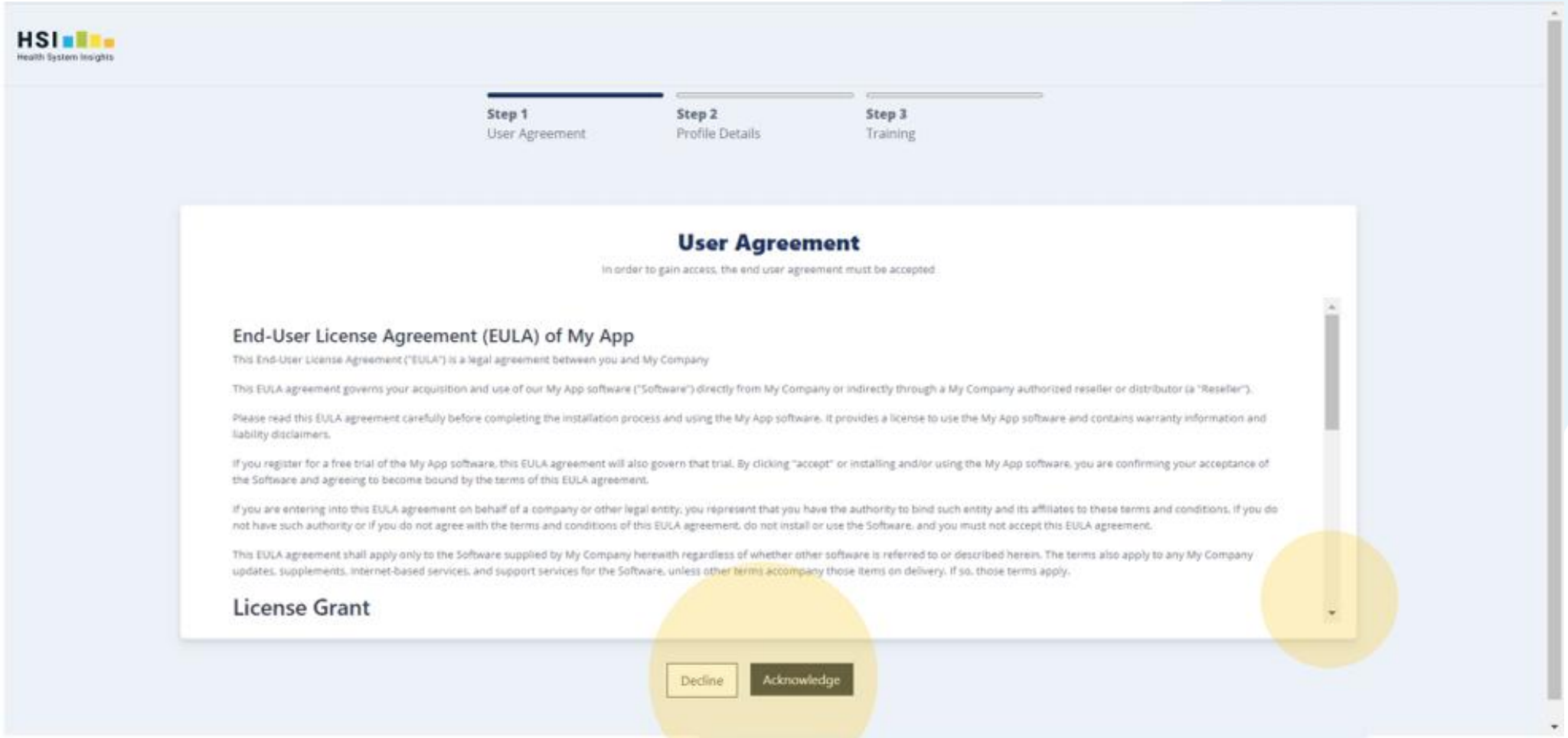
Navigate to https://hsi.ontariohealth.ca and click the **ONE® ID** button



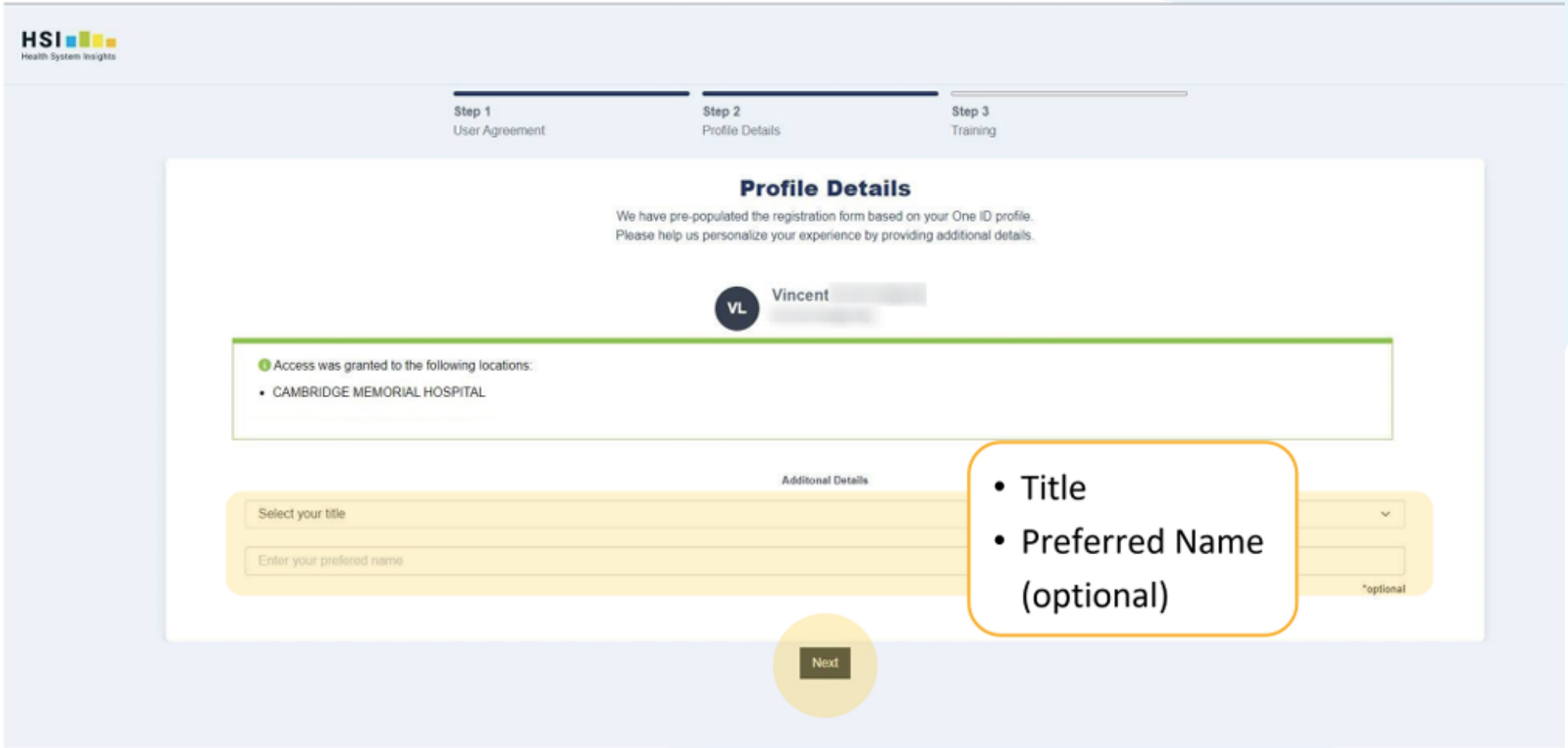
Log in with your ONE® ID Credentials



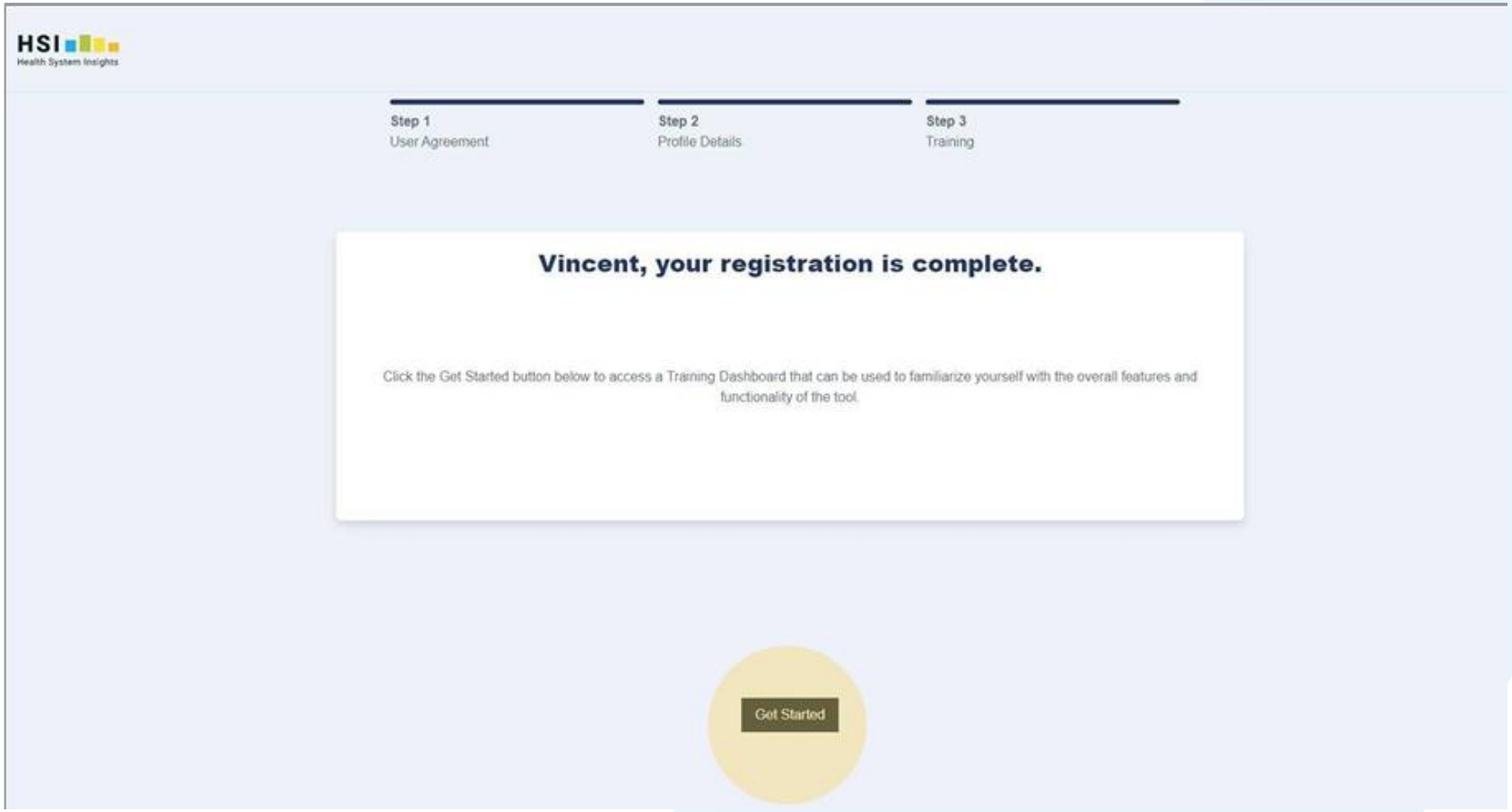
Accept the End User License Agreement



Complete Your Profile Details



Acknowledgement



Note: Access to the dashboard will remain in a pending status until your facility's HSI FRA approves your request.

RT Data and HSI

- RT Reports Dashboard

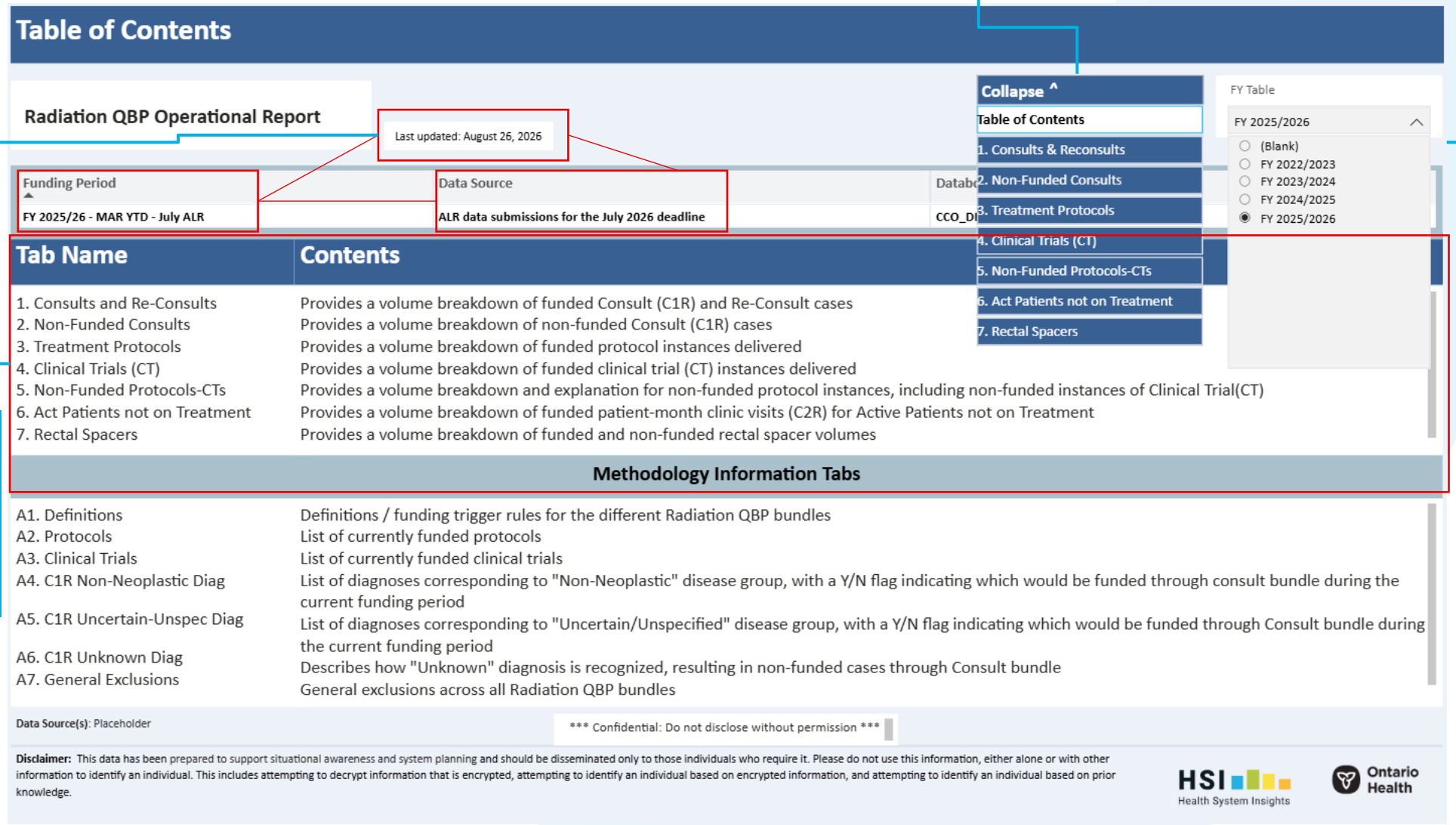
Navigating the Dashboard

This section provides information on where the data is coming from and when it was last updated

This section provides an overview of each tab and what data is included

Use this drop-down menu to navigate through different pages

Use this drop-down menu to filter for the fiscal year



Consults & Reconsults

This tab provides a breakdown by Disease Group of funded Consult (C1R) and Re-Consult case volumes at the facility.

See sections "Consult Definition" and "Re-Consult Definition" in Appendix "A1. Definitions" for more detail about funding trigger rules for the Consult and Re-Consult bundles.

Navigation Tab

FY Table

FY 2025/2026

Funded Consults and Re-Consults - Case Volumes

Disease Group	Sum of Consults	Sum of Reconsults
BREAST	1391	41
GENITOURINARY	1163	35
GASTROINTESTINAL	751	50
LUNG	701	87
HEAD AND NECK	686	40
HAEMATOLOGY	523	14
SKIN	466	20
GYNAECOLOGICAL	443	27
SARCOMA	428	30
CENTRAL NERVOUS SYSTEM	235	7
ENDOCRINE	186	10
BENIGN NEOPLASMS	175	0
UNCERTAIN/ UNSPECIFIED SITES	74	2
NON-NEOPLASTIC DIAGNOSES	61	0
PRIMARY UNKNOWN	46	1
Total	7372	366

This table provides an overview of the Funded Consults and Re-Consults volumes by disease group.

Data Source(s): Placeholder

*** Confidential: Do not disclose without permission ***

Disclaimer: This data has been prepared to support situational awareness and system planning and should be disseminated only to those individuals who require it. Please do not use this information, either alone or with other information to identify an individual. This includes attempting to decrypt information that is encrypted, attempting to identify an individual based on encrypted information, and attempting to identify an individual based on prior knowledge.

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Non-Funded Consults

This tab provides a volume breakdown for non-funded consult volumes, which are defined as non-malignant. This may be the result of a data quality issue or a result of clinical practice issues. Facilities should review patient level data to understand whether the consults listed are the result of a data quality or clinical practice issue and resubmit data as required.

See section "Consult Definition" in Appendix "A1. Definitions" for more detail about funding trigger rules for the Consult bundle.

Navigation Tab

FY Table

FY 2025/2026

Diagnoses	Count of YTD Volume	% of Total
NON-NEOPLASTIC DIAGNOSES	37	0.5%
UNCERTAIN/ UNSPECIFIED SITES		
Total	37	0.5%

Sum of Consults	Total %
7372	99.5%

Total C1R Volumes
7409

This tab shows the Non-Funded Consults, including the Unknown and Uncertain Volumes.

Unknown and Non-Neoplastic Volumes
(by ICD-10-CA code)

Diagnoses	Diagnoses Description	Count of YTD Volume
NON-NEOPLASTIC DIAGNOSES	Amyotrophic lateral sclerosis	1
NON-NEOPLASTIC DIAGNOSES	Anaemia unspecified	1
NON-NEOPLASTIC DIAGNOSES	Aneurysm & dissectn othr spec art	1
NON-NEOPLASTIC DIAGNOSES	Ankylosis of joint lower leg	7
NON-NEOPLASTIC DIAGNOSES	Disease of gallbladder unspecified	1
NON-NEOPLASTIC DIAGNOSES	Disease of salivary gland unspecified	1
NON-NEOPLASTIC DIAGNOSES	Disorder of breast unspecified	2
Total		37

Uncertain/Unspecified Volumes
(by ICD-O-3 morphology and ICD-10-CA topography codes)

Diagnoses	Diagnoses Description	Count of YTD Volume

Data Source(s): Placeholder

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Treatment Protocols

This tab provides a volume breakdown of funded radiation protocol instances delivered at the fa ivered to treat a primary cancer are listed under "Primary Protocol Volumes", wheras those delivered to t Volumes".

The Treatment bundle is triggered when an evidence-informed protocol is delivered to a patient rotocol having been completed during the current funding period. See section "Protocol Definition" in A ng trigger rules.

These two tables outline the Primary and Metastatic protocols and their volumes across the disease groups.

Navigation Tab

FY Table

FY 2025/2026

Primary Protocols

Disease Group(Note that '*' refers to any disease)

All

Disease Group(Note that '*' refers to any disease)	Protocol	Funding Band	Number of Instances
BREAST	BREAST_CF	Band 1	594
BREAST	BREAST_FULL_CF	Band 1	594
BREAST	BREAST_FULL_HYPO	Band 1	594
BREAST	BREAST_FULL_SHORT_COURSE	Band 1	594
BREAST	BREAST_HYPO	Band 1	594
BREAST	BREAST_NODAL_ONLY_CF	Band 1	594
BREAST	BREAST_NODAL_ONLY_HYPO	Band 1	594
BREAST	BREAST_PAR_CF	Band 1	594
BREAST	BREAST_PAR_HYPO	Band 1	594
BREAST	BREAST_PAR_UHYPO	Band 1	594
Total			4884

Metastatic Protocols

Protocol

All

Disease Group(Note that '*' refers to any disease)	Funding Band	Protocol	Number of Instances
*	Band 1	BONE_MET_CON_MULTI	2132
*	Band 2	BONE_MET_CON_MULTI	884
*	Band 3	BONE_MET_CON_MULTI	16
CNS	Band 2	BONE_MET_CON_MULTI	7
CNS (PEDS)	Band 4	BONE_MET_CON_MULTI	2
GYNECOLOGY	Band 2	BONE_MET_CON_MULTI	2
HEMATOLOGY	Band 1	BONE_MET_CON_MULTI	111
HEMATOLOGY	Band 2	BONE_MET_CON_MULTI	41
LUNG	Band 2	BONE_MET_CON_MULTI	4
SARCOMA	Band 3	BONE_MET_CON_MULTI	36
Total			3276

Data Source(s): Placeholder

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Clinical Trials

This tab provides a volume breakdown of funded radiation Clinical Trials delivered at the facility during the current funding period. Clinical Trials delivered to treat a primary cancer are listed under "Primary Intent", wheras those delivered to treat metastases are listed under "Metastatic Intent".

The Treatment bundle is triggered when a Clinical Trial (CT) is delivered to a patient, with the last radiation course of treatment for this CT having been completed during the current funding period. See section "Clinical Trial Definition" in Appendix "A1. Definitions" for more detail about funding trigger rules.

Navigation Tab

FY Table

FY 2025/2026

Primary Protocols

Disease Group	Clinical Trail	Funding Band	Number of Instances
GENITOURINARY	NCT06409910	Band 3	30
GENITOURINARY	NCT05328505	Band 3	12
GENITOURINARY	NCT06235697(PEL)	Band 2	12
BREAST	NCT05592938	Band 1	11
GENITOURINARY	NCT06235697(PROS)	Band 2	5
BREAST	NCT04228991	Band 3	4
GENITOURINARY	NCT00913939	Band 5	1
GENITOURINARY	NCT05364229	Band 2	1
Total			76

Metastatic Protocols

Disease Group	Clinical Trial	Funding Band	Number of Instances
*	NCT05717166	Band 2	1
Total			1

These two tables outline the Clinical Trials for the Primary and Metastatic protocols.

Data Source(s): Placeholder

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Non-Funded Protocols - CTs

This tab provides a volume breakdown for Non-funded protocol instances, including non-funded instances of Clinical Trials (CT). This may be the result of a data quality issue or a result of clinical practice issues. Facilities should review patient level data to understand whether the protocol instances listed are the result of a data quality or clinical practice issue and resubmit data as required.

See sections "Protocol Definition" and "Clinical Trial Definition" in Appendix "A1. Definitions" for more detail about funding trigger rules for the Treatment bundle.

Navigation Tab

FY Table

FY 2025/2026

Instances	Sum of NFP Instances	% of Total Non-Funded Protocol Instances
Gyne_Brachy Auditing	3	0.0%
Imaging Unresolved (applies to non-pediatric Metastatic Protocols only) within 30 days		
Invalid Protocol Name submitted		
Miscoded Metastatic Protocol Intent		
Total	11	0.1%

Sum of Funded Protocol Instances	% of Total Funded Protocol Instances
8237	99.9%

Total Protocol Instances Submitted
8248

- Miscoded Primary Protocol Intent or Unknown Intent (Intent "unknown" or not coded with the expected "Primary" intent)
- Miscoded Metastatic Protocol Intent (Intent not coded with the expected "Metastatic" intent)
- Miscoded or Unknown Disease (Disease Group is unknown or does not correspond to protocol specified)
- Invalid Protocol Name Submitted (Protocol name is missing or is not recognized according to protocols/clinical trials listed on tabs "A1. Protocols" and "A2. Clinical Trials")
- Miscoded Ready to Treat (RTT) Date (RTT date should be the same for all consecutive courses representing the same protocol/clinical trial)
- Imaging "Unresolved" (Metastatic protocol instance not associated with any imaging activity, be it MR or CT scan)

RT Protocol	Sum of NFP Instances

This page outlines the Non-Funded Protocol Instances. You can select each of the categories on the left to get more details on the protocols.

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Active Patients not on Treatment (C2R Visit Volumes) (Patient-Months)

This tab provides a volume breakdown by Disease Group of funded patient-month clinic visits (C2R) at the facility for Active Patients not on Treatment

See section "Active Patients not on Treatment Definition" in Appendix "A1. Definitions" for more detail about funding trigger rules for this bundle.

Navigation Tab

FY Table

FY 2025/2026

Active Patients not on Treatment - C2R Visit Volumes
(Patient-Months)

Disease Group	Sum of Number of Patients (ALR Cases)	Sum of Number of C2R-Patient-Months
GENITOURINARY	717	1198
BREAST	369	506
HEAD AND NECK	262	397
ENDOCRINE	252	363
GASTROINTESTINAL	181	295
GYNAECOLOGICAL	138	218
LUNG	122	217
SARCOMA	127	173
SKIN	117	166
HAEMATOLOGY	110	147
BENIGN NEOPLASMS	82	113
CENTRAL NERVOUS SYSTEM	50	69
UNCERTAIN/ UNSPECIFIED	50	65
Total	2631	3996

This table indicates the volumes for Active Patients not on Treatment by Disease Group.

Data Source(s): Placeholder

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Rectal Spacers

This tab provides a volume breakdown of funded rectal spacers delivered at the facility during the current funding period. Rectal spacer indications are listed "Rectal Spacer Description" as well as corresponding NHPIP activity code.

Rectal Spacer Funding is triggered when a patient with prostate cancer is treated with a primary intent prostate evidence-informed protocol with the following indications (Pts with IBD, Pts on chronic anti-coagulant therapy, Pts with bilat Hip Prosthesis, RC SAP Approved) and reported rectal spacer insertion (399) is completed during the current funding period.

See section "Rectal Spacer" in Appendix "A1. Definitions" for more detail about funding trigger rules.

Navigation Tab

FY Table

FY 2025/2026

Funded Rectal Spacer Volumes

Description	NHPIP	Sum of Value
Pts on chronic anti-coagulant therapy	727x	35
RC SAP Approved	729x	14
Pts with IBD	726x	4
Pts with bilat Hip Prosthesis	728x	3
Total		56

Non-Funded Rectal Spacers

Non-Funded Rectal Spacers	Total
NHPIP_399	12

These two tables provide an overview on both Funded and Non-Funded Rectal Spacers.

Data Source(s): Placeholder

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HSI Health System Insights

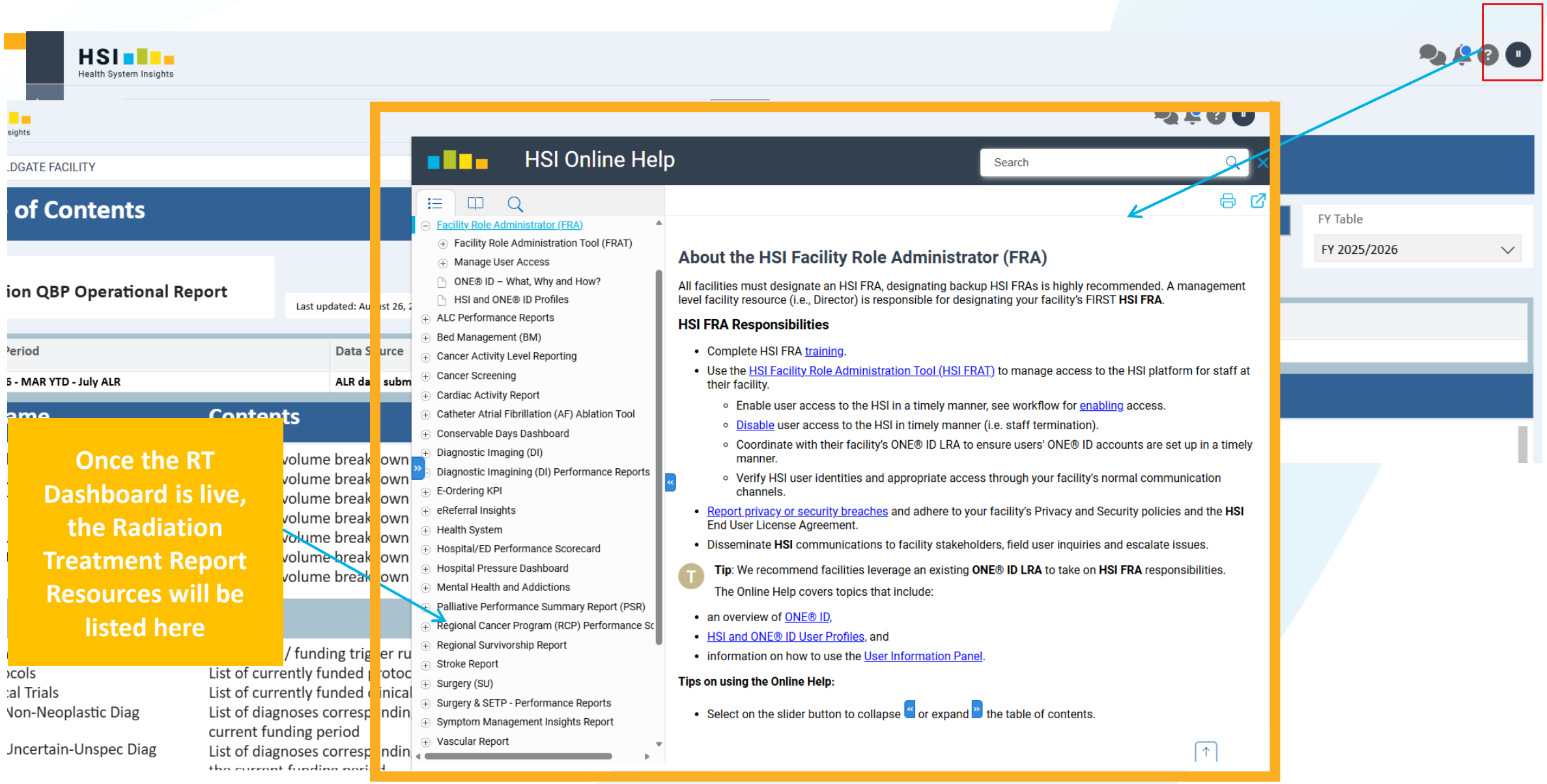
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Resources

HSI Online Help

Access by clicking on ‘How to Register’ on log in screen or the ? icon once logged in




HSI
Health System Insights
of Contents
ion QBP Operational Report
Last updated:
Period
Data Source
- MAR YTD - July ALR
ALR data submitted
Contents
volume breakdown
List of currently funded protocols
List of currently funded clinical Trials
List of diagnoses corresponding current funding period
Non-Neoplastic Diag
Uncertain-Unspec Diag
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Once the RT Dashboard is live, the Radiation Treatment Report Resources will be listed here
HSI Online Help
Search
Facility Role Administrator (FRA)
Facility Role Administration Tool (FRAT)
Manage User Access
ONE® ID – What, Why and How?
HSI and ONE® ID Profiles
ALC Performance Reports
Bed Management (BM)
Cancer Activity Level Reporting
Cancer Screening
Cardiac Activity Report
Catheter Atrial Fibrillation (AF) Ablation Tool
Conservable Days Dashboard
Diagnostic Imaging (DI)
Diagnostic Imagining (DI) Performance Reports
E-Ordering KPI
eReferral Insights
Health System
Hospital/ED Performance Scorecard
Hospital Pressure Dashboard
Mental Health and Addictions
Palliative Performance Summary Report (PSR)
Regional Cancer Program (RCP) Performance Sc
Regional Survivorship Report
Stroke Report
Surgery (SU)
Surgery & SETP - Performance Reports
Symptom Management Insights Report
Vascular Report
About the HSI Facility Role Administrator (FRA)
All facilities must designate an HSI FRA, designating backup HSI FRAs is highly recommended. A management level facility resource (i.e., Director) is responsible for designating your facility's FIRST HSI FRA.
HSI FRA Responsibilities
Complete HSI FRA training.
Use the HSI Facility Role Administration Tool (HSI FRAT) to manage access to the HSI platform for staff at their facility.
Enable user access to the HSI in a timely manner, see workflow for enabling access.
Disable user access to the HSI in timely manner (i.e. staff termination).
Coordinate with their facility's ONE® ID LRA to ensure users' ONE® ID accounts are set up in a timely manner.
Verify HSI user identities and appropriate access through your facility's normal communication channels.
Report privacy or security breaches and adhere to your facility's Privacy and Security policies and the HSI End User License Agreement.
Disseminate HSI communications to facility stakeholders, field user inquiries and escalate issues.
Tip: We recommend facilities leverage an existing ONE® ID LRA to take on HSI FRA responsibilities.
The Online Help covers topics that include:
an overview of ONE® ID,
HSI and ONE® ID User Profiles, and
information on how to use the User Information Panel.
Tips on using the Online Help:
Select on the slider button to collapse or expand the table of contents.

Need More Help?

If you have any further questions, please reach out to:
QBPPowerBIsupport@ontariohealth.ca for any **PowerBI dashboard related inquiries**