A Registrars Defense to ACoS Accreditations

NAACCR 2018-2019 WEBINAR SERIES

NAACCR

Q&A

Please submit all questions concerning the webinar content through the Q&A panel.

If you have participants watching this webinar at your site, please collect their names and emails

We will be distributing a Q&A document in about one week. This document will fully answer questions asked during the webinar and will contain any corrections that we may discover after the webinar.

NAACCR

Guest Speaker

Sara Morel, CTR

• Lead Network CTR/Tumor Registry. MidMichigan Health

Courtney Jagneaux, RHIA, CTR

Senior Registry Operations Director, RegistryPartners

NAACCR

Agenda

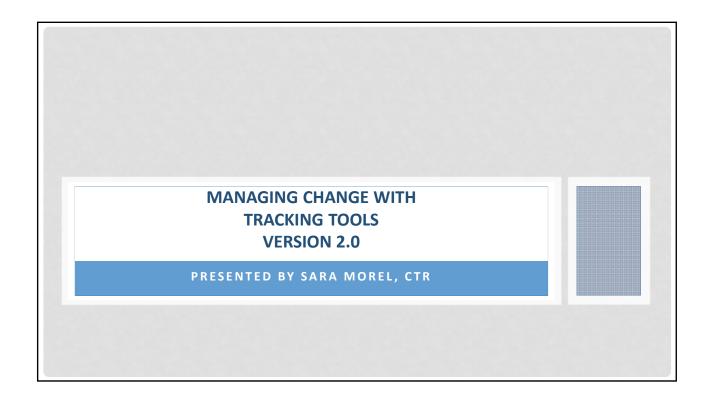
Managing Change with Tracking Tools Version 2.0

•Sara Morel

Survival Guide to Survey Preparations & NCDB Tools

Courtney Jagneaux

NAACCR



OBJECTIVES

Version 2.0: Managing change in 2019

- Provide updated standards tracking templates to be able to prepare for an Commission on Cancer Survey with a focus on Integrated Network Cancer Programs.
- Introduce new tracking and reporting templates for the Rectal Cancer Center for Excellence Standards for the NAPRC Accreditation
- Review EPIC Case Finding reporting that provides improved workflows.
- No relevant disclosures

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TOPICS TO BE COVERED

- NAPRC Tracking templates (National Accreditation Program for Rectal Cancer)
- NAPBC Breast Tracking (National Accreditation Program for Breast Cancer for Cancer Conferences)
- Cancer Conference Reporting
- Commission on Cancer Standards Tracking
- EPIC Reporting
- Integrated Network Cancer Program Cancer Committee Minutes Template
- Survey Prep Templates
- CTR Education Tracking Template
- Updated Abstracting Guide

NAPRC: CHAPTER 1 RC-MDT: RECTAL CANCER MULTIDISCIPLINARY TEAM

- NAPRC Standard 1.2: RC-MDT: The rectal MDT team must have at least 1 appointed physician
 from each specialty: Pathology, Radiology, Surgery, Medical and Radiation Oncology. All
 surgeons who are performing rectal cancer surgeries must be required members of the RCMDT-Rectal Cancer Multidisciplinary Team. This must be documented in the rectal MDT
 meeting minutes. The team also must appoint a Rectal Cancer Program Director and a Rectal
 Cancer Program Coordinator.
- NAPRC Standard 1.2: Alternates: Alternates may be appointed annually but this must be
 done at the first meeting of the year. This must be documented in the rectal leadership
 meeting minutes. Surgeons will not have alternates due to that they are all required to be on
 this MDT team and no required member is allowed to be an alternate for more than one role.
- NAPRC Standard 1.2: Meetings: The standard recommends that these RC-MDT/Rectal Cancer
 Conferences be held at a different times from other cancer sites cancer conferences;
 however if all specialties are present they may be held in conjunction with a general or other
 site cancer conference. These are the meetings where the standards will be discussed and
 documented.

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NAPRC: RC-MDT RECTAL CANCER MULTIDISCIPLINARY TEAM ATTENDANCE

- NAPRC Standard 1.3: Required Attendance Percentage: Each required member/alternate
 appointed is required to attend at least 50% and this is monitored by the Rectal Cancer Program
 Director. If you need to appoint more than 2 groups for each specialty this is allowed but each
 grouping must meet the 50% attendance requirement.
- NAPRC Standard 1.3: Discussion and Participation MDT/Rectal Cancer Conference: Video
 Conferencing is allowed as long as any member has access to review all radiology and pathology
 slides and that they can participate in the RC MDT/Cancer Conference. Each rectal cancer case
 must be presented twice at cancer conferences; pre and post op.
- NAPRC Standard 1.4: Frequency of RC-MDT/Rectal Cancer Conference: Must meet at least twice
 per month and it is required for at least 1 physician member from each specialty to attend each
 RC-MDT/Cancer Conference.
- NAPRC Standard 1.5: RC MDT/Cancer Conference Minutes: Each year the 2 monthly meetings/cancer conferences must be documented along with documented compliance with the required standards.

NAPRC: RC-MDT CANCER CONFERENCE ATTENDANCE TEMPLATE RECTAL CANCER CONFERENCE (MDT) ATTENDANCE 2019 CORE GROUP 2019 (Can miss 12 total for the entire year) Date Date Date Surgery Pathology Radiology Medical Oncology Radiation Oncology Rectal Cancer Program Director Rectal Cancer Coordinator Tumor Registry vellow=absent CORE GROUP 2019 Date Date Date Surgery Pathology Radiology Medical Oncology Radiation Oncology Rectal Cancer Program Director Rectal Cancer Coordinator Tumor Registry x=present 11

NAPRC: RECTAL CANCER PROGRAM DIRECTOR AND RECTAL CANCER PROGRAM COORDINATOR

- NAPRC Standard 1.1: CoC Accreditation: Before moving forward with an NAPRC Accreditation the facility must be Accredited with the Commission on Cancer.
- NAPRC Standard 1.5: Rectal Cancer Program Director is responsible to:
 - · Chair the Rectal MDT/Cancer Conferences
 - Will be the liaison between the NAPRC and CoC. Must attend at least 1 CoC meeting annually to present the rectal data once available.
 - Must review and present the NCDB Data (at least 4 times per year)
 - Must do a chart review for the Chapter 2 Standards
- NAPRC Standard 1.6: Rectal Cancer Program Coordinator is responsible to:
 - Provides support for the MDT/Cancer Conferences
 - Monitor and coordinate the care for rectal cancer patients from diagnosis through treatment with policies and procedures in place.
- NAPRC Standard 1.7: Rectal Cancer Program Required Education
 - Required education modules for surgeons, pathologists and radiologists of the RC MDT will be assigned annually and completed certificates must be uploaded annually.

NAPRC: CHAPTER 2: PATHOLOGY AND STAGING

NAPRC Standard 2.1: Review of Diagnostic Pathology

- 95% of all new rectal cancer patients must have their biopsy confirmed by an appointed Pathologist before the start of any treatment. If done outside the report/slides must be reviewed and documented in the medical record.
 - Example: Our pathologists review all slides and reports for the Rectal Cancer Conferences so
 we document this review in our Tumor Board Note in EPIC.
 - If the pathology slides are not available a RC MDT report must be reviewed and documented by an RC MDT pathologist.
 - If neither are available then the patient must be re-biopsied.
 - 20% of cases will be randomly audited at the survey (max 100 cases)

NAPRC Standard 2.2: Staging to be completed before definitive treatment

- 95% of all new rectal cancer cases must be staged before the start of definitive treatment and must be documented in the medical record.
- Systemic and Local staging
 - CT Scan or PET Scan of the Chest, Abdomen and Pelvis & MRI of the Rectum (a combined CT/PET is acceptable however if a PET is done without a CT, it does not meet this standard)
 - 20% of cases will be randomly audited at the survey (max 100 cases)
 - · Use of EPIC staging forms.

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NAPRC: CHAPTER 2 MRI AND CEA

• NAPRC Standard 2.3: MRI Reporting (Standardized)

- 90% of new rectal cancer cases who have not had treatment yet must have an MRI of the rectum performed and the report must be read by an appointed radiologist of the RC MDT.
- 95% of new rectal cancer cases who have not had treatment, the MRI reports must be standardized and must contain all required elements for staging.
- 20% of cases will be randomly audited at the survey (max 100 cases)

NAPRC Standard 2.4: CEA Labs

- For 75% of new rectal cancer cases who have not had treatment yet; they must have a CEA-Carcinoembryonic Antigen level drawn and this must be documented in the medical record.
- 20% of cases will be randomly audited at the survey (max 100 cases)

NAPRC: CHAPTER 2 RECTAL PLANNING MDT DISCUSSION (CANCER CONFERENCE) NAPRC Standard 2.5: Treatment Planning Discussion (Must be done for 100% of cases; 20% review, max 100 cases) 1st time being presented at cancer conference template of the required items for the Agenda: • Pre Op Information: (1st time presented) • DOB, age & sex: • Site: RECTUM • MRN: · Clinical Diagnosis: Presenting physician/navigator: · Other physicians: Pathology: · Question for the pathologist: Clinical AJCC stage:CT Chest/ABD/Pelvis imaging date: PET imaging date: MRI imaging date: · Reason for review: · Colonoscopy outcomes: Pre-Treatment CEA: Pre-Treatment MSI: Additional Information: • Discussion included: Clinical Trial, Genetics, Palliative Care, Social Services, Rehab or Plastic Surgery · Referrals to: Medical Oncology, Radiation Oncology Intent for treatment: Curative, Control, Maintenance, Palliative, Supportive Tumor Board note is done in EPIC as well 15

NAPRC: CHAPTER 2 TREATMENT EVALUATION LETTER TO PRIMARY CARE PHYSICIAN

- NAPRC Standard 2.6: Treatment Evaluation/Discussion Summary (from Cancer Conference/Tumor Board)
 - This must be completed for at least 50% of all rectal cancer patients.
 - 20% of cases will be randomly audited at the survey (max 100 cases)
- Letter to PCP with recommendations from Cancer Conference Template
 - · Patient Name, Date of Birth, MRN
 - Dear Dr
 - Example: Items included:
 - Thanking them for their referral
 - · Date patient was presented at a Rectal Cancer Conference
 - · Presenting physician and specialties in attendance
 - We include what was presented for example: clinical history and any pertinent imaging and pathology.
 - NCCN Guidelines and recommendations for treatment
 - · Clinical trials status if eligible
 - The presenting physician signs off on note in EPIC and this goes to the PCP/referring physician.
 - All of this gets documented in the Oncology History in EPIC by the Rectal Cancer Coordinator.

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NAPRC: CHAPTER 2 DEFINITIVE TREATMENT, SURGICAL RESECTION AND PATHOLOGY

- NAPRC Standard 2.7: Definitive treatment (to start within 60 days from initial clinical eval-bx/dx)
 - 80% are required to have 1st course treatment within 60 days
 - 20% of cases will be randomly audited at the survey (max 100 cases)
- NAPRC Standard 2.8: Surgical Resection
 - 80% of cases are required to have surgery by an appointed Rectal Surgeon on the MDT
 - 95% of operative notes are required to be in standardized synoptic format
 - 20% of cases will be randomly audited at the survey (max 100 cases)
- NAPRC Standard 2.9: Pathology CAP Reports
 - 90% of case are to be read by an appointed pathologist on the MDT
 - 95% of cases are required to be in the CAP template and reported out within 2 weeks from surgery
 - 20% of cases will be randomly audited at the survey (max 100 cases)
- NAPRC Standard 2.10: Photographs of Surgical Specimens
 - 65% of case are to be required to be photographed—anterior, posterior and lateral views (must be presented at cancer conference)
 - 20% of cases will be randomly audited at the survey (max 100 cases)

NAPRC: CHAPTER 2 TREATMENT OUTCOMES DISCUSSION AT **CANCER CONFERENCE**

- Standard 2.11: Treatment Outcomes Discussion (Template for Cancer Conference/Tumor Board Agenda)
- Post Op information: (2nd time presented) (20% review, max 100 cases)
 - Imaging: None requested unless otherwise specified
 - Patient name, DOB, age & sex:
 - Site: RECTUM
 - MRN:
 - Presenting physician/navigator:
 - Final Pathological Diagnosis & Final Pathological AJCC Stage:
 - Prior date presented at Cancer Conference:
 - Neo-Adj treatment before surgery:
 - Neo-Adj treatment date of completion:

 - Date of surgery and Type of surgery:
 - Surgical approach:
 - Presence or absence of stoma:
 - Post-Op complications:
 - Unexpected findings:
 - Specimen photographs:
 - Tumor Location:
 - Indication of sphincter involvement, CRM margin status & distal margin status:
 - Tumor Regression Grade & Mesorectal Grade:
 - Discussion included: Clinical Trial, Genetics, Palliative Care, Social Services, Rehab or Plastic Surgery
 - Referrals to: Medical Oncology, Radiation Oncology, Palliative Care, Nutrition, PT, Ostomy
 - Tumor Board note is done in EPIC as well

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NAPRC: CHAPTER 2 TREATMENT OUTCOME SUMMARY SURVIVORSHIP CARE PLAN

- Standard 2.12: Treatment Outcome Summary/SCP
 - 50% of eligible surgical patients must have a treatment summary provided within 4 weeks of surgery. A copy is provided to the PCP/referring physician.
 - 20% of cases will be randomly audited at the survey (max 100 cases)
- Standard 2.13: Adjuvant Treatment after the Surgical Resection
 - 50% of surgery cases that are eligible should begin any adjuvant treatment within 8 weeks from their definitive surgery of the primary rectal cancer tumor. This referral process and monitoring of patients is done by the Rectal Cancer Coordinator.
 - 20% of cases will be randomly audited at the survey (max 100 cases)
- Chapter 3 in progress: RQRS, NCDB data and measures
- Policies and Procedures

| | NAPRC: REG | CTAL | STAN | DARE | os co | MPLIANCE | |
|-------------------------------------|--|------------------------------|----------------|---------------------------|---------------------------|--|----|
| Review of the st | tandards and chapter 2 char | t audits n | nust be d | ocument | ed annual | ly | |
| Tracking templa | | | | | | | |
| | • | Nationa | l Rectal Cance | Accreditation | Program | | |
| | | | | | | | |
| | | | | | | | |
| | Standards | MDT meeting discussed at: | | MDT meeting discussed at: | MDT meeting discussed at: | Outcome/Action Items from last date approved | |
| | Standard 1.1 Commission on Cancer | | | | | | |
| | Accreditation Standard 1.2 Rectal Cancer Multidisciplinary | | | | | | |
| | Care | | | | | | |
| | Standard 1.3 Rectal Cancer Multidisciplinary Team Attendance | | | | | | |
| | Standard 1.4 Rectal Cancer Multidisciplinary Team Meetings | | | | | | |
| | Standard 1.5 Rectal Cancer Program Director | | | | | | |
| | Standard 1.6 Rectal Cancer Program Coordinator | | | | | | |
| | Standard 1.7 Rectal Cancer Program Education | | | | | | |
| | Standard 2.1 Review of Diagnostic Pathology | | | | | | |
| | Standard 2.2 Staging Before Definitive Treatment | | | | | | |
| | | | | | | | 21 |

NAPRC: PATIENT TRACKING AND CHAPTER 2 CHART AUDIT Patient name and Date of Birth: Facility and MRN: Date Standards Notes Date of biopsy, facility & pathologist Biopsy reviewed by a MDT pathologist Date clinicals taging done, physician and stage If previous ly diagnosed is the clinical/pathological staging noted in EPIC? 1-Standard 2.1 2-Standard 2.2 EPIC? Imaging done (CT CAP or PET s can) Date MRI done, facility, physician Is the MRI report in the synoptic reporting format? Date Pre-Treatment CEA, facility and level Date initially presented at cancer conference and all items discussed before definitive treatment 3-Standard 2.2 4-Standard 2.3 5-Standard 2.4 Date noted in EPIC that the patient was presented (Tumor Board Note) Date treatment evaluation and recommendations created (Letterto PCP-Communication 7-Standard 2.6 Date patient began 1st course definitive treatment, facility, physician Total number of days from tissue biopsyto first treatment (goal 60 8-Standard 2.7 Date patient began 1st cours e definitive treatment, facility, physician Total number of days from tissue biapsyto first treatment (goal 60 days) Date of surgical resection 8, physician Date pathology report date, CAP and Ostrich compliant Date surgery photos presented at cancer conference and pathologist: (must be within 4 weeks—28 days) Days from surgery to 2nd presentation at cancer conference 2nd date presented at cancer conference post up and physician: Tumor Boar Obis cussion nated in EPIC 9-Standard 2.8 10-Standard 2.9 11-Standard 2.10 12-Standard 2.11 Tumor Board Discussion noted in EPIC Date Treatment Summary/Survivorship CP due date (due within 4 weeks from date in 2.11) Date adjuvant treatment started (within 8 weeks) from surgery from 13-Standard 2.12 22

CANCER CONFERENCE AGENDA TEMPLATE Example for General CoC CoC: Standard 1.7 · Patient name: • DOB, age & sex: • Site: • MRN: • Diagnosis: · Presenting: Other physicians: Pathology: • Stage: Imaging: Reason for Review: · Additional Information or questions for pathology or radiology: • Discussion included: Clinical Trial, Genetics, Palliative Care, Social Services, Rehab or Plastic Surgery 23

CANCER CONFERENCE AGENDA TEMPLATE Example for Breast NAPBC: Standard 1.2 Patient Name: • DOB, Age & Sex: • MRN: • BMI: Presenting and Other physicians: Site: Diagnosis, Grade, ER/PR, HER2, KI67: Stage: • Imaging & Pathology: Surgery type and date: Genetics eligible or Clinical Trials eligible: · Chief Complaint: Prior Mammogram & Past Medical and Surgical History: · Signs and Symptoms: Smoking and Alcohol History: · Family History of Cancer: Menopause Status: • Discussion included: Palliative Care, Social Services, Rehab, Plastic Surgery · Treatment Plan: • Intent for treatment: Curative, Control, Maintenance, Palliative, Supportive 24

PREPARING CANCER CONFERENCE AGENDA'S

Workflow for Preparing Cancer Conference Agenda's

- Create Word Agenda using template
- InBasket name "Cancer Conference"
- Create a new InBasket Message
- · Distribution List in EPIC and Managing it
- Options-Set expiration date for message
- Subject line-labeling per facility
- Copy and paste from the Word document into the body of the message and click Accept
- Frequency of sending each agenda
- · Accessibility to physicians and staff attending
- Ability to add cases through EPIC messaging

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AFTER CANCER CONFERENCE REPORTING

UDF:

- CC NCCN Guidelines Followed
- CC Clinical Trials Discussed
- CC Genetics Discussed
- CC Palliative Care Discussed
- CC Psychosocial Discussed
- CC Rehab Discussed
- CC Plastic Surgery Discussed
- CC Discussion of workup/stage
- CC Prognostic Factors Discussed
- CC Last name presenting physician (text)

General Page:

- Presentation CA Conf & Date of CA Conf
- Query Wizard Report
- Patients presented multiple times, Total case presentations & Overall tracking

CANCER REGISTRY QUALITY CONTROL STANDARD 1.6 QA REVIEWER TEMPLATE

CoC Standard 1.6

- Patient name & Date of birth:
- Facility & MRN:
- Primary site & laterality if applicable:
- Histology:
- · Class of case:
- Date of 1st contact:
- Date of initial diagnosis:
- Date of 1st course treatment: (CTR to list treatment):
- · Clinical grade:
- Pathological grade:
- Post treatment grade:
- Clinical TNM and Stage group:
- · Pathological TNM and Stage group:
- · Evidence of this cancer:
- Date of 1st recurrence if applicable: NA (unless known recurrence):
- Case finding source & Abstracted by:
- Date reviewed: _____ Initials:
- Abstracting timeliness: All cases are abstracted within 3-6 months from date of diagnosis or date of first contact.
 All unknown fields or fields coded as 9 are reviewed on a monthly basis for all cases.
- All cases are submitted when requested to the NCDB.

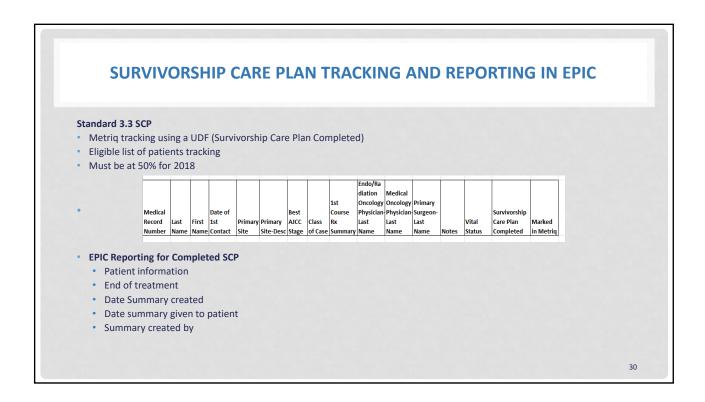
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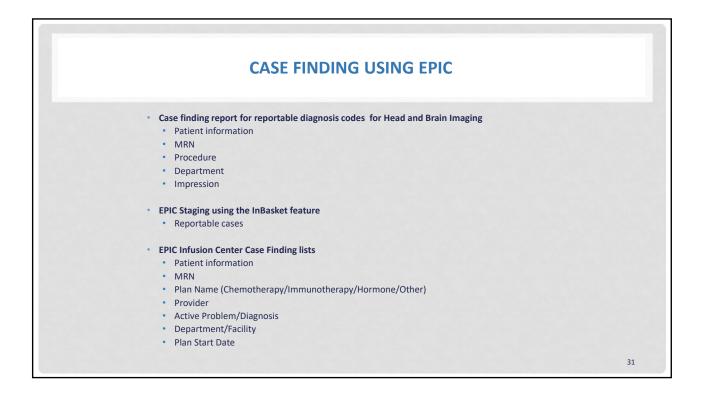
DISTRESS TRACKING STANDARD 3.2

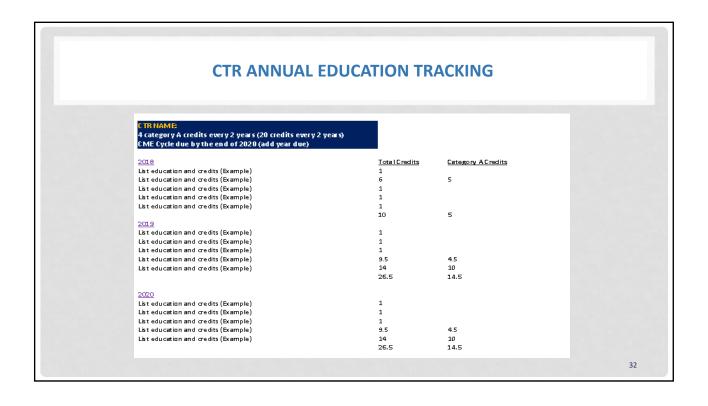
CoC: Standard 3.2 Distress Scores

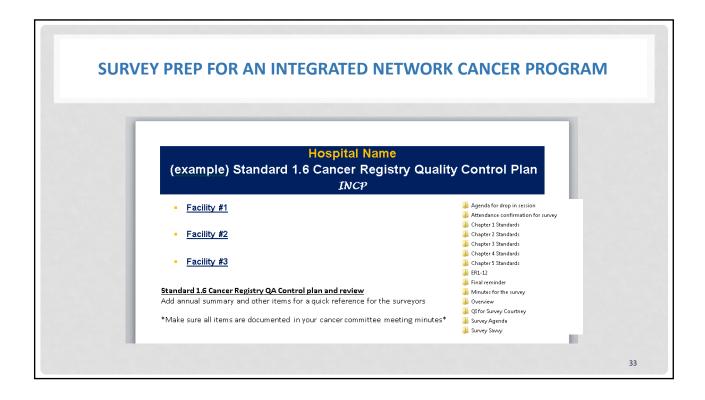
- · Distress routing in EPIC to Tumor Registry InBasket
- Quarterly report for Social Workers/Nurses
- Metriq UDF (Psych Distress Screen Score)
- Query Wizard report

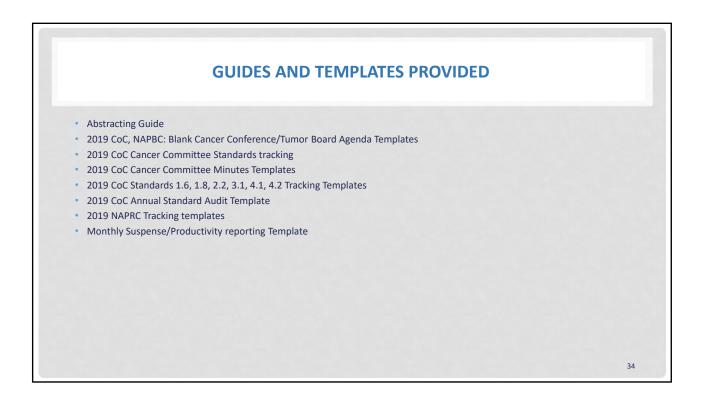
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| Medical Oncology | | | | | | | | | | | | | | | | | | | | |
| Radiation Oncology | | | | | | | | | | | | | | | | | | | | |
| Surgery | | | | | | | | | | | | | | | | | | | | |
| Pathology | | | | | | | | | | | | | | | | | | | | |
| Radiology | | | | | | | | | | | | | | | | | | | | |
| Yellow-absent | | | | | | | | | | | | | | | | | | | | |
| x=present | | | | | | | | | | | | | | | | | | | | |
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| Radiology | | | | | | | | | | | | | | | | | | | | |
| Yellow-absent | | | | | _ | | | | _ | | | | _ | _ | | | | _ | | |
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| - Present | | | | | | | | | | | | | | | | | | | | |
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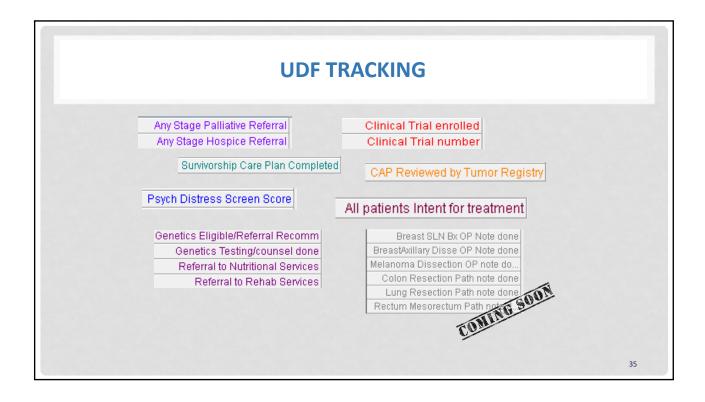


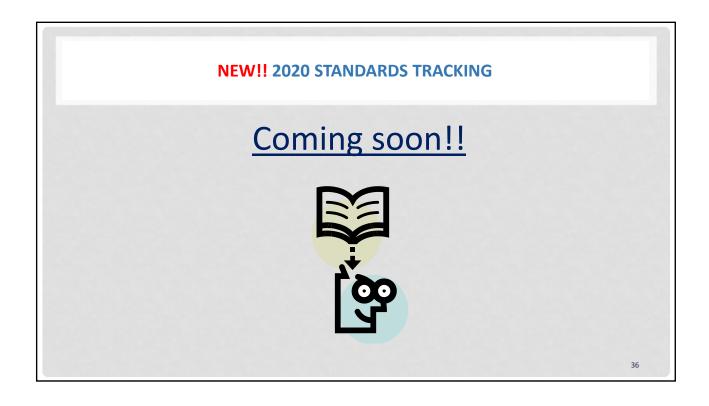






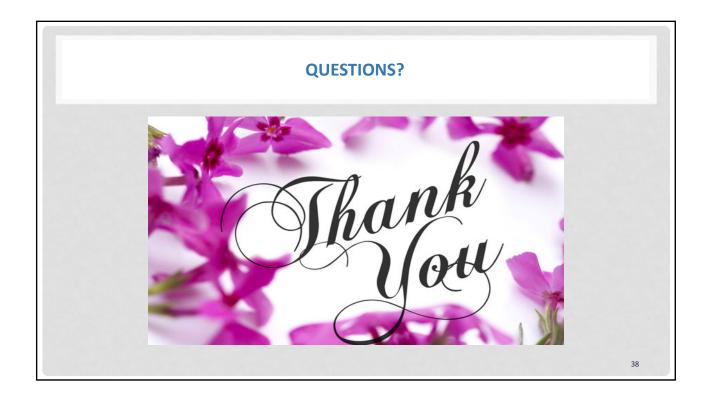


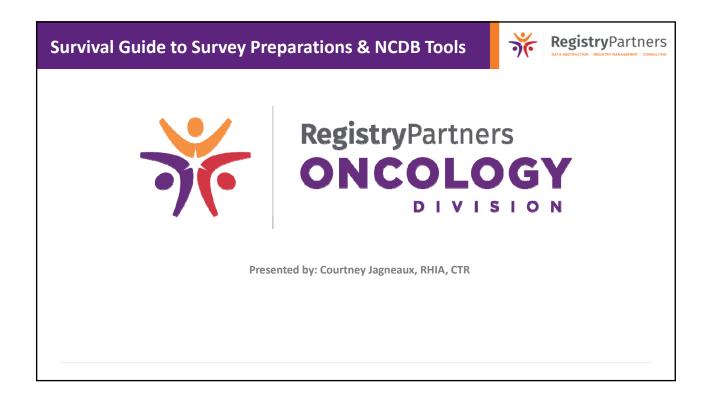




REFERENCES

- Commission on Cancer Program Standards; https://www.facs.org/quality-programs/cancer/coc/standards
- Breast NAPBC Standards: https://www.facs.org/quality-programs/napbc/standards
- Rectal NAPRC Standards: https://www.facs.org/quality-programs/cancer/naprc/standards
- Abstracting Guide: AJCC Staging Manual, Collaborative Staging-prior to 2018, ICD-O, SEER Coding Manuals, Hematopoietic Database, SEER RX, Multiple Primaries Manual, MCSP Manual, STORE manual, SEER Website, NAACCR Website.
- Thanks to Wendy Johnson, CTR, Ginger Greenwood, CTR, Maggie Nelson, CTR, and Tara Talaski CTR for assisting with reviewing and editing these presentation slides.





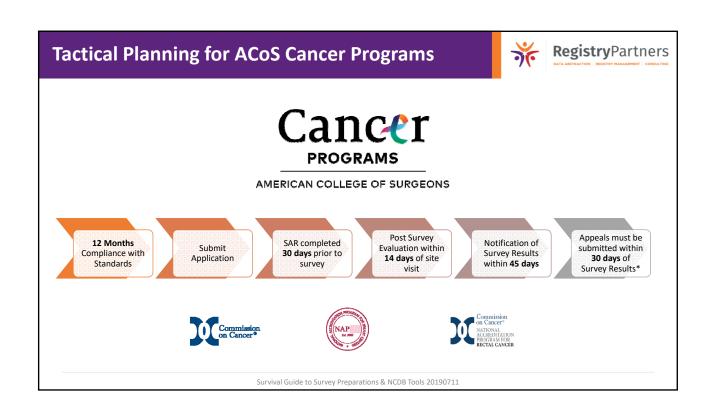


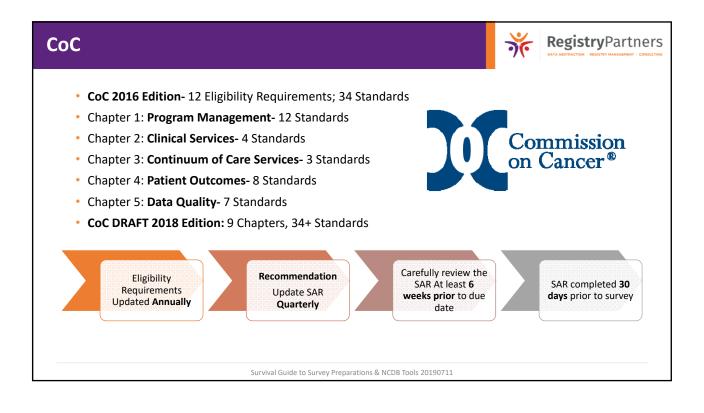
Learning Objectives

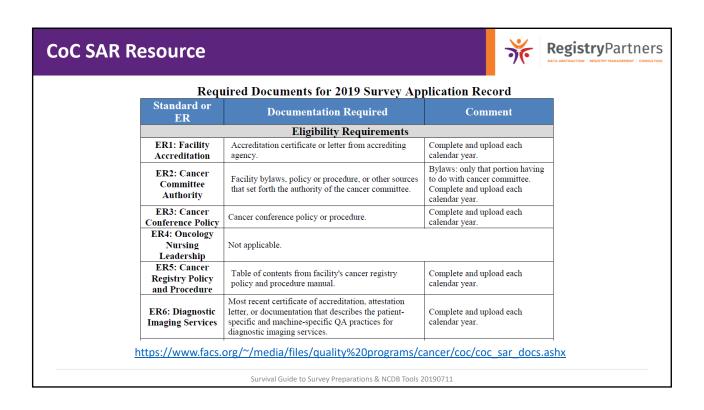


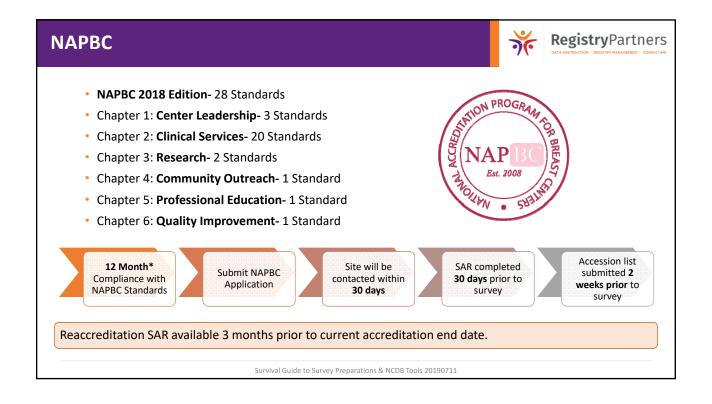
- Identify best practices for preparing a cancer program for survey
- Provide tools to assist in survey application documentation
- Educate registrars on how to support their Cancer Liaison Physician
- Review NCBD Reporting tools and best practices

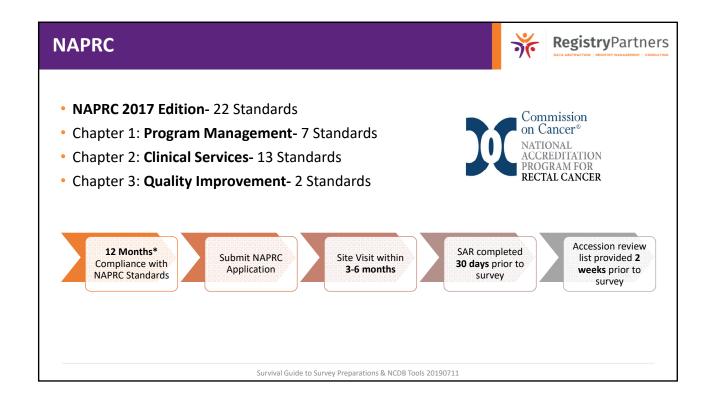












Accreditation Preparation





Survival Guide to Survey Preparations & NCDB Tools 20190711

Survival Guide to Survey Preparations





✓ Planning

• Set your program up for success in advance with best practice tools

√ Timeline

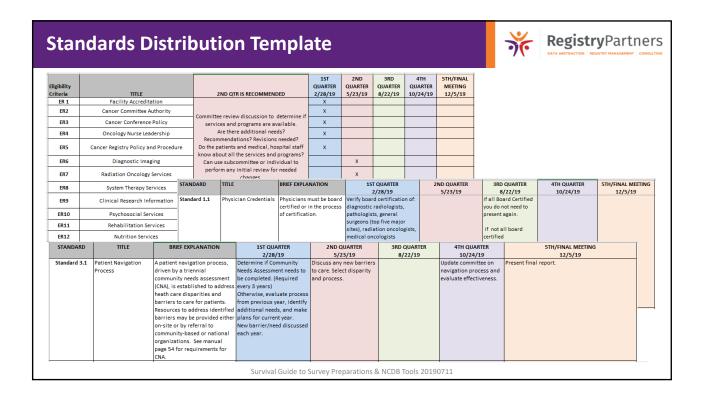
• Communicate schedules and deadlines to your committee members

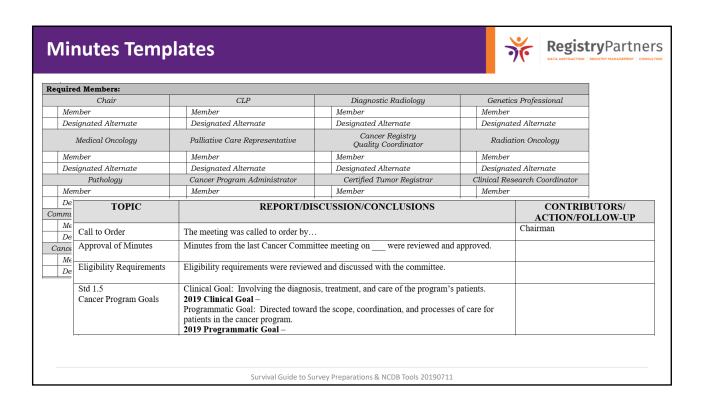
✓ Organization

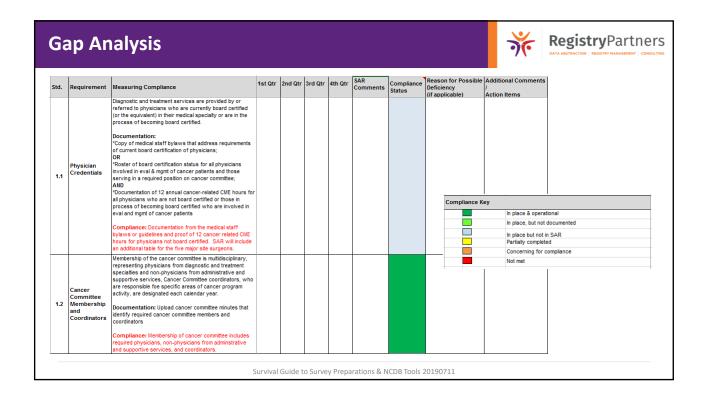
• Establish a system that allows for multiple contributors

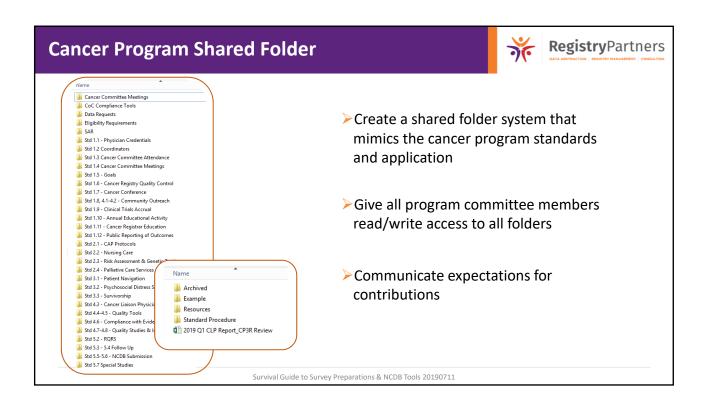
✓ Team Work

· Hold all cancer committee members accountable for their roles and responsibilities









Supporting Your Cancer Liaison Physician



Understanding the CLP Role

- <u>Primary Responsibilities</u>: monitor, interpret, and provide update reports of the program's performance using NCDB data to evaluate and improve the quality of care.
- Secondary Responsibilities:
 - Report on CoC activities, initiatives, and priorities to the cancer committee.
 - Serve as liaison between the cancer program, the CoC, and the American Cancer Society.
 - · Attend the CoC on-site survey and meets with the surveyor

Being "In the Know"

- Complete CLP orientation within three months of initial appointment AND on reappointment every three years.
- Know the ACS Representative & State Chair
- Attend CLP Breakfast sessions when possible
- Subscribe to the NewsCLiPs
- Subscribe to the CoC Newsbreaks
- Using NCDB Reporting Tools



https://www.facs.org/quality-programs/cancer/clp

Survival Guide to Survey Preparations & NCDB Tools 20190711

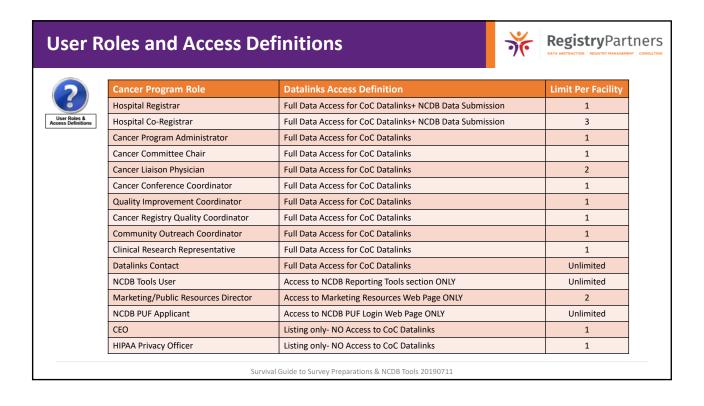
NCDB Tools

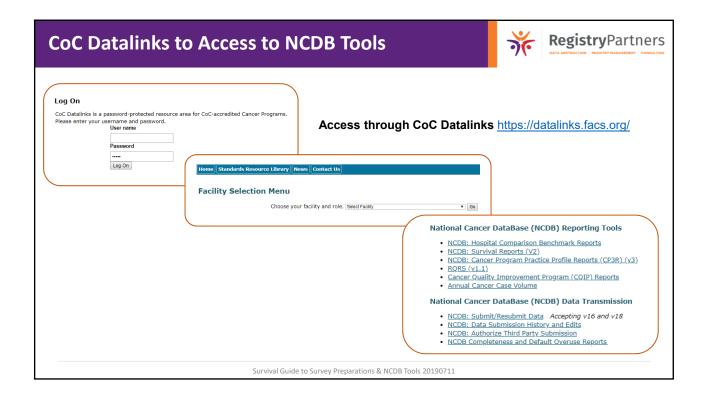


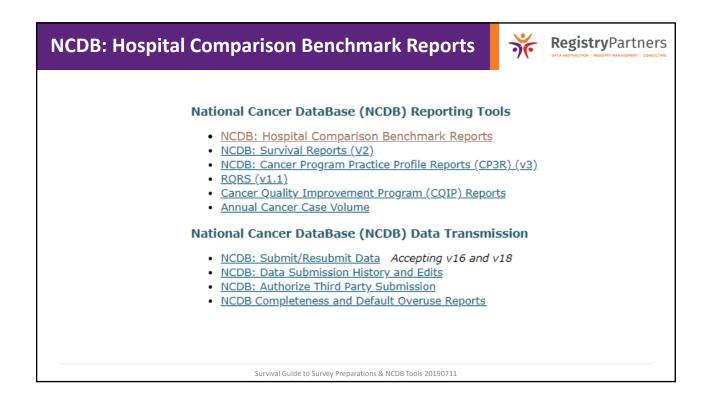


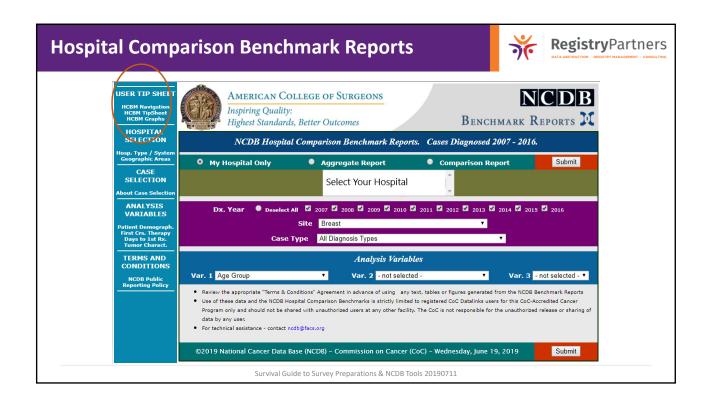
- o NCDB Hospital Comparison Benchmark Reports (HCBR)
- NCDB Survival Reports (Survival)
- o Cancer Program Practice Profile Reports (CP3R)
- Rapid Quality Reporting System (RQRS)
- Cancer Quality Improvement Program (CQIP)
- o Annual Cancer Case Volume
- o Data Completeness and Default Overuse Report

https://www.facs.org/quality-programs/cancer/ncdb/qualitytools









Hospital Comparison Benchmark Report



- My Hospital Only
 - Create comparisons of data reported to the NCDB from your own cancer registry
 - Assess up to 3 variables

Aggregate Report

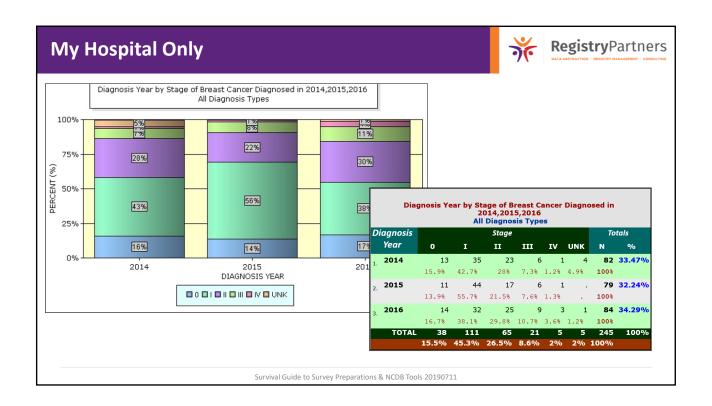
- · Create reports made from combined data at the hospital system, state, region, or at the national level.
- Assess up to 3 variables

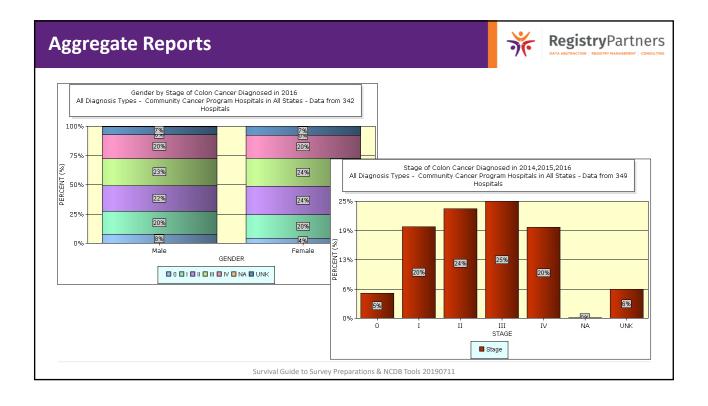
Comparison Report

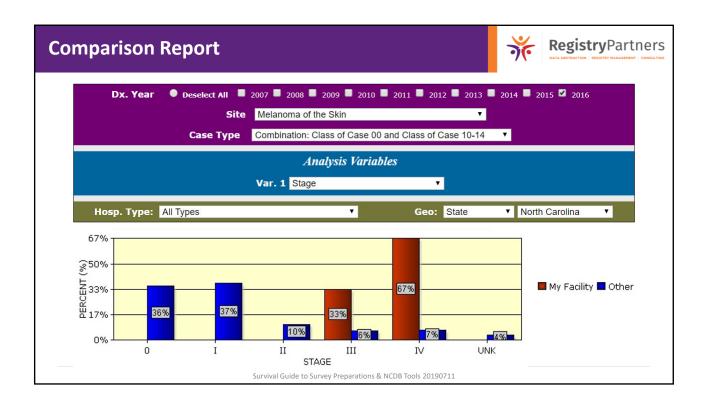
- Create comparison reports of the cases submitted to the NCDB by your cancer program and all the other programs identified in your comparative group
- Assess only 1 variable

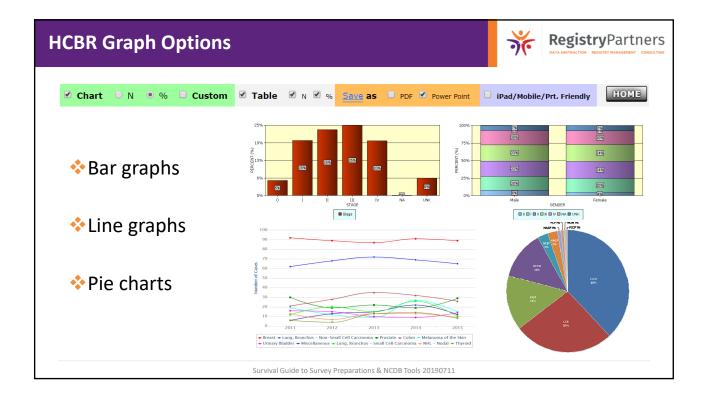
Benchmark Reports Analysis Variables

| Age Group | Histology | Zip Code Level Income |
|------------------|------------------------------|-----------------------|
| Stage | Tumor Behavior | Days to Treatment |
| Diagnosis Year | First Course Treatment | First Course Surgery |
| Gender | Charleston Comorbidity Score | Radiation Therapy |
| Race/Ethnicity | Distance Traveled | Systemic Therapy |
| Insurance Status | Zip Code Level Education | <u> </u> |

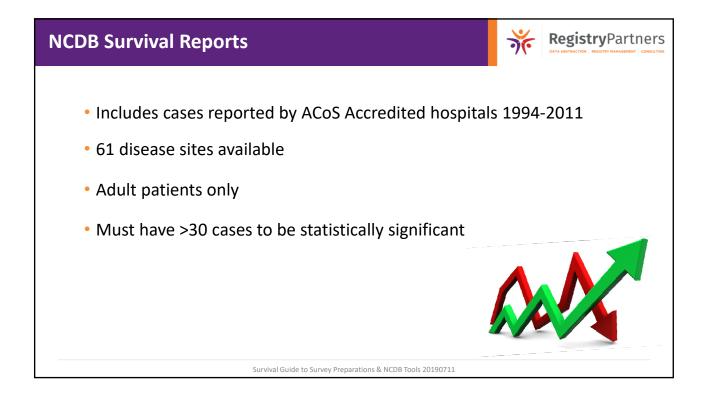


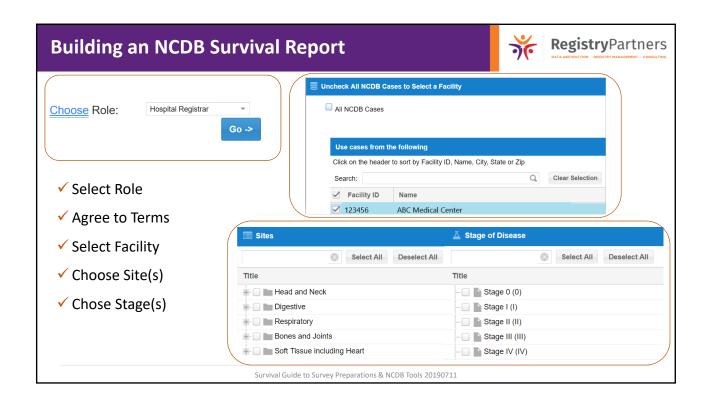


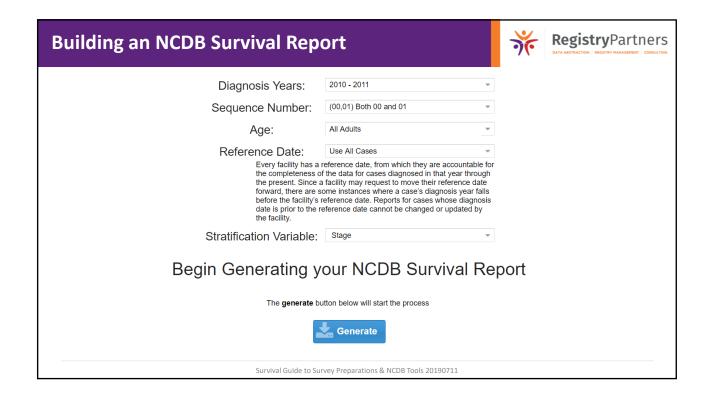


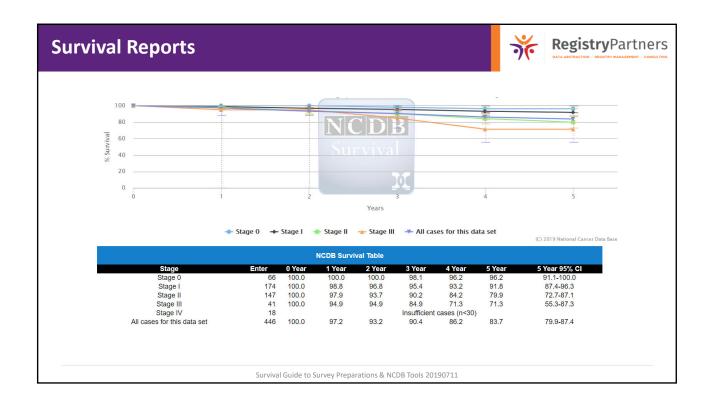


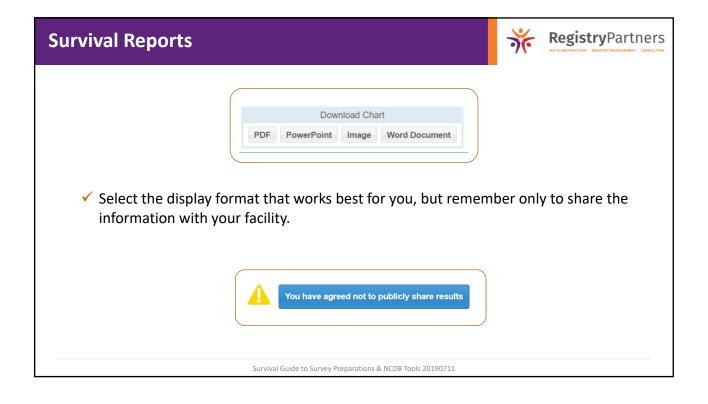


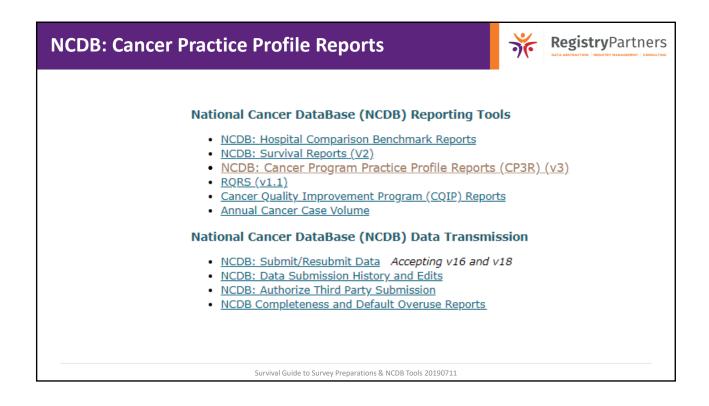


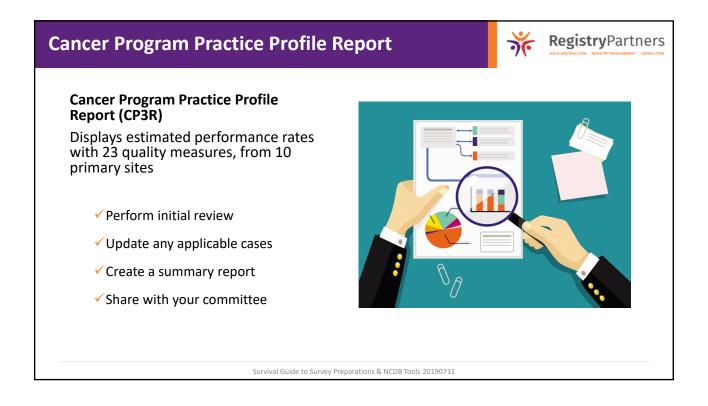


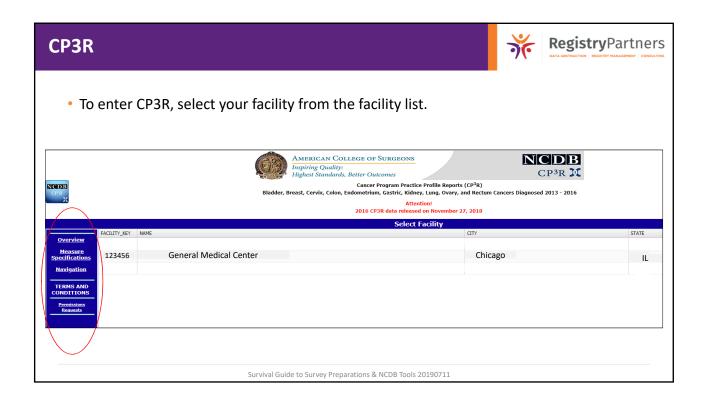


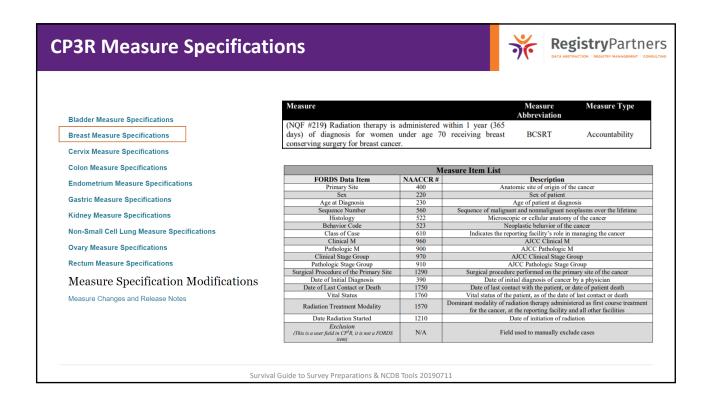














Standards 4.4 & 4.5 – Measure Type, Definition & Use



Accountability (Std 4.4)

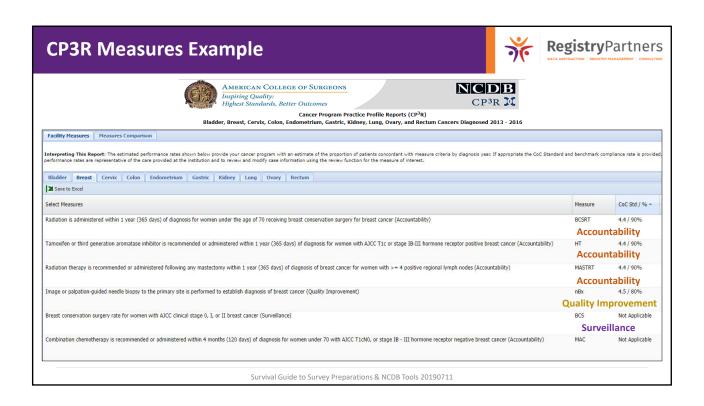
High level of evidence supports the measure, including multiple randomized control trials.
 These measures can be used for public reporting, payment incentive programs, and the selection of providers by consumers, health plans, or purchasers

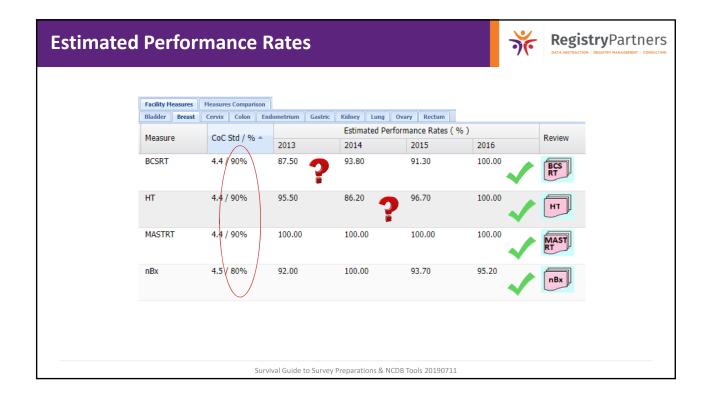
Quality Improvement (Std 4.5)

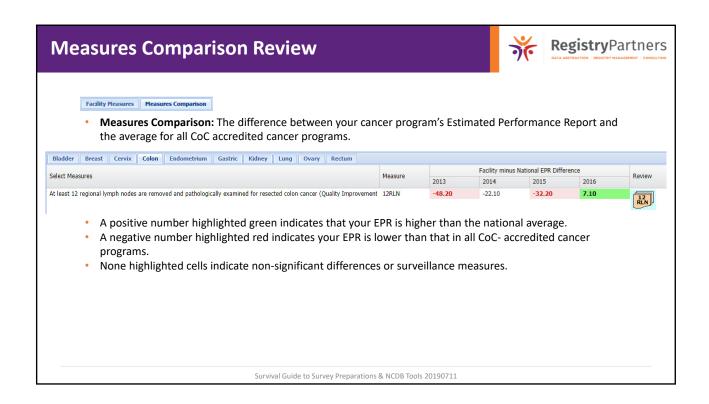
• Evidence from experimental studies, not randomized clinical trials support the measure. Intended for internal monitoring of performance within an organization

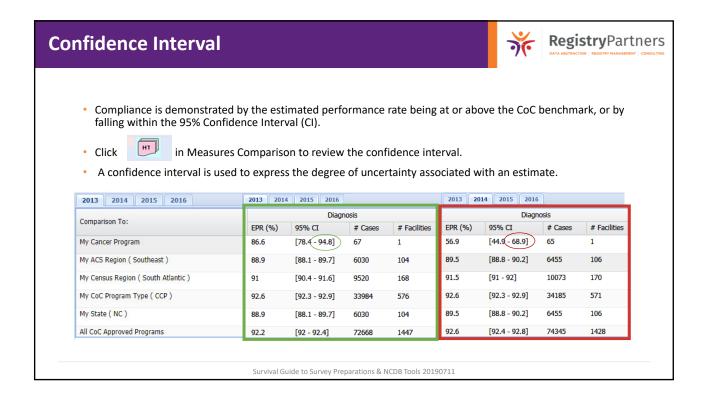
Surveillance

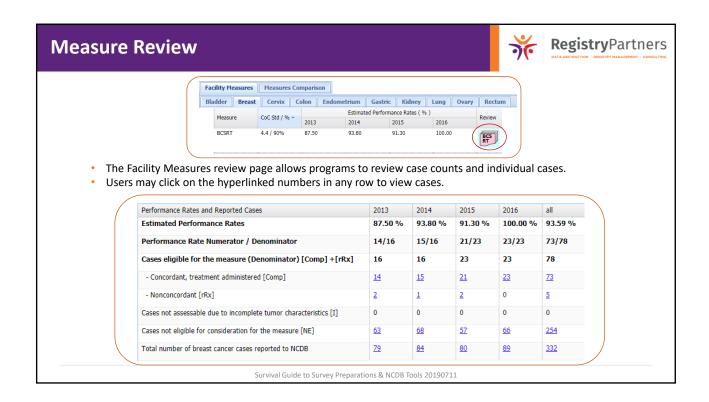
 Limited evidence exist that supports the measure or the measure is used for informative purposes to accredited programs. These measures can be used to identify the status quo as well as monitor patterns and trends of care in order to guide decision-making and resource allocation

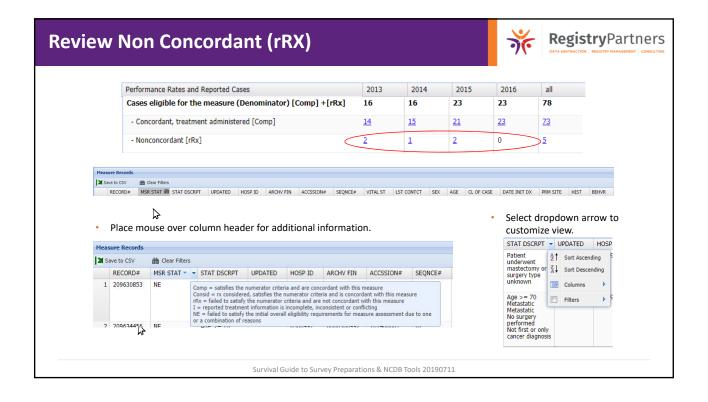


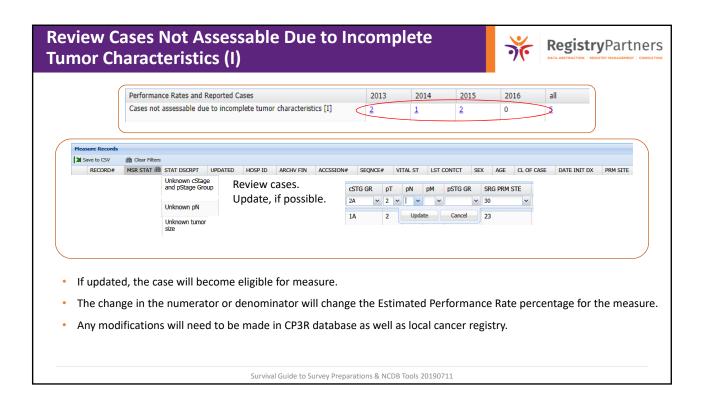


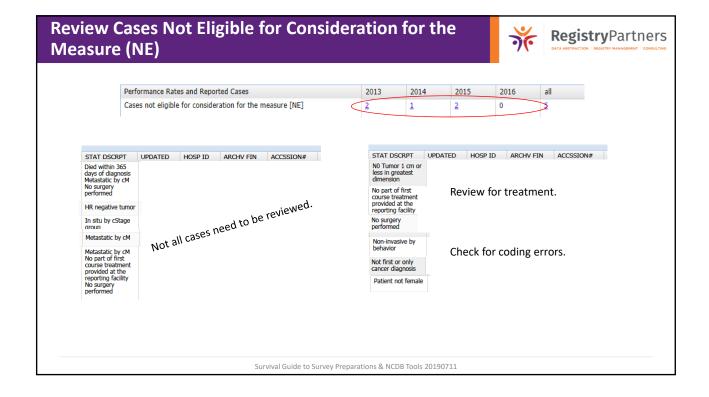


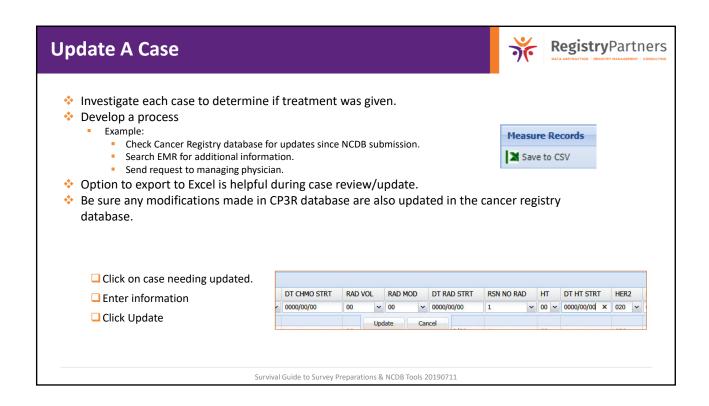












CP3R Presentation to Cancer Committee



A Summary is presented by the Cancer Liaison Physician (CLP) at least once per year.

| Quality Measure | EPR | PR | 95% CI | Cases | Action Plan/Comments |
|--|-------------------|-------|--------------|-------|---|
| BREAST [HT]: Tamoxifen or third generation aromatase inhibitor is considered or administered within 1 year (365 days) of diagnosis for women with AJCC T1c or stage IB-III hormone receptor positive breast cancer. Std 4.4 | 90% | 53.8 | (41.7, 61.9) | 35/65 | Non-compliant. 28 cases-no documentation of HT Given. 2 cases-HT started >365 days. |
| BREAST [nBx]: Image or palpation-guided needle biopsy (core or FNA) is performed to establish diagnosis of breast cancer Std 4.5 | 80% | 100% | (100, 100) | 57/57 | Compliant. No Action Needed |
| BREAST [BCS]: Breast conservation surgery rate for women with AJCC clinical stage 0, I, or II breast cancer. (Surveillance) | Not Applicable | 57.4% | (43.3, 71.5) | 27/47 | No EPR. Not applicable. No Action Needed |
| CERVIX [CBRRT]: Use of brachytherapy in patients treated with primary radiation with curative intent in any stage of cervical cancer. (Surveillance) | Not Applicable | 100% | (100, 100) | 2/2 | No EPR. Not applicable. No Action Needed |
| COLON [12RLN]: At least 12 regional lymph nodes are removed and examined for resected colon cancer. Std 4.5 | 85% | 69.2% | (44.1, 94.3) | 9/13 | Compliant. No Action Needed. |

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CP3R Action Plan



An action plan is developed and executed if programs performance rates are below the CoC's EPR

| Quality Measure | EPR | PR | 95% CI | Cases | Action Plan/Comments |
|--|-----|-------|--------------|-------|--|
| BREAST [HT]: Tamoxifen or third generation aromotase inhibitor is considered or administered within 1 year (365 days) of diagnosis for women with AICC T1c or stage IB-III hormone receptor positive breast cancer. Std 4.4 | 90% | 85.7% | (72.7, 98.7) | 24/28 | Compliant. No Action Needed |
| | | 53.8 | (41.7, 61.9) | 35/65 | 28 cases-no documentation of HT Given. 2 cases-HT started >365 days. |

Quality Measure: HT – Tamoxifen or third generation aromatase inhibitor is considered or administered within 1 year (365 days) of diagnosis for women AJCC T1c or Stage 1B-Stage 3 hormone receptor positive breast cancer.

Expected Performance Rate: 90% Actual Performance Rate: 53.8%

Action Plan: Reviewed 28 cases with no information and found that managing physicians were from the same physician group that will not respond to our request for treatment information. The Cancer Program administrator and CLP agreed to meet with the administrator from the physician group. They will explain the importance of the information and the impact it has on our cancer program. They will request electronic access to the physician group's patients.

Effectiveness: The physician group agreed to give us access for 30 days to the patients that needed additional treatment information. The Cancer Registry will submitted a list of patients and update once access is granted.

RQRS



National Cancer DataBase (NCDB) Reporting Tools

- NCDB: Hospital Comparison Benchmark Reports
- NCDB: Survival Reports (V2)
- NCDB: Cancer Program Practice Profile Reports (CP3R) (v3)
- RQRS (v1.1)
- Cancer Quality Improvement Program (CQIP) Reports
- · Annual Cancer Case Volume

National Cancer DataBase (NCDB) Data Transmission

- NCDB: Submit/Resubmit Data Accepting v16 and v18
- NCDB: Data Submission History and Edits
- NCDB: Authorize Third Party Submission
- NCDB Completeness and Default Overuse Reports

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RQRS- Minimum Requirements

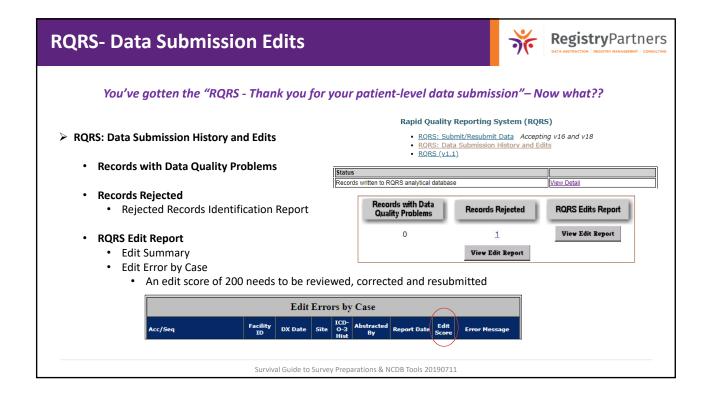


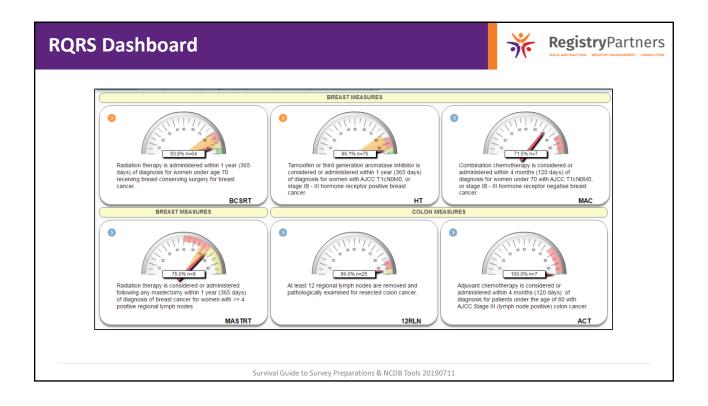
- NCDB needs basic information from a pathology report, an assigned accession number, and a sequence number to assess cases for RQRS adjuvant therapy measures.
- Records may be submitted at any time and in varying states of completeness.
 - To be compliant with Standard 5.2, records must be submitted quarterly.
 - For commendation of Standard 5.2, records must be submitted monthly.
 - I recommend to submit weekly or bi-monthly depending on your program size.
- The minimum data requirements are found in Table 1. Table 2 documents all required data element for the measure algorithms to run. (page 9)

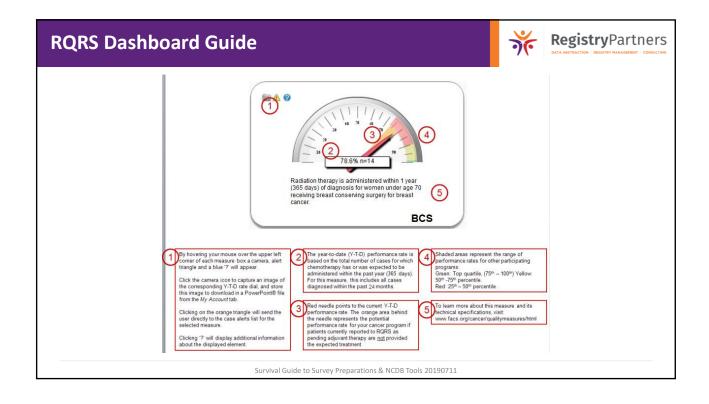
Rapid Quality Reporting System (RQRS)

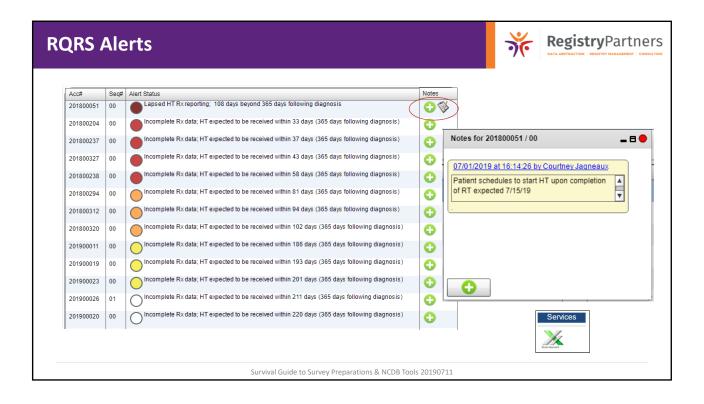
- RQRS: Submit/Resubmit Data
- RQRS: Data Submission History and Edits
- RQRS (v1.1)

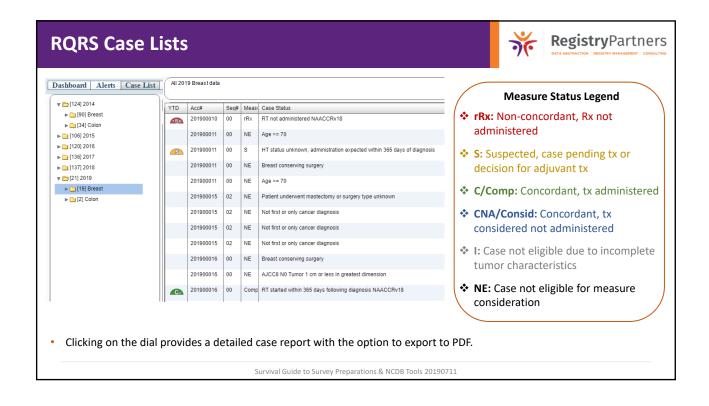
https://www.facs.org/~/media/files/quality%20programs/cancer/ncdb/rgrs_userguide.ashx

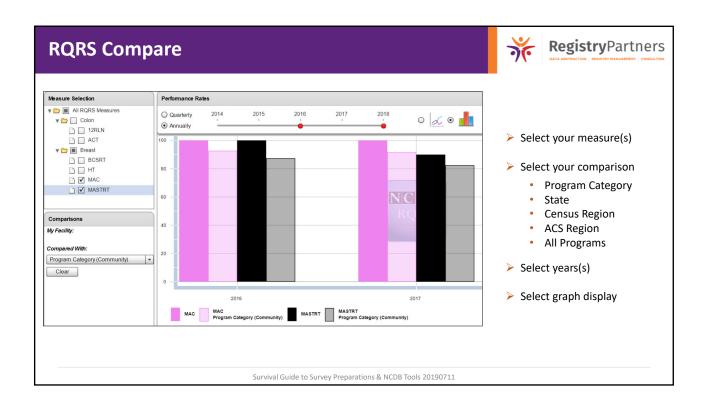


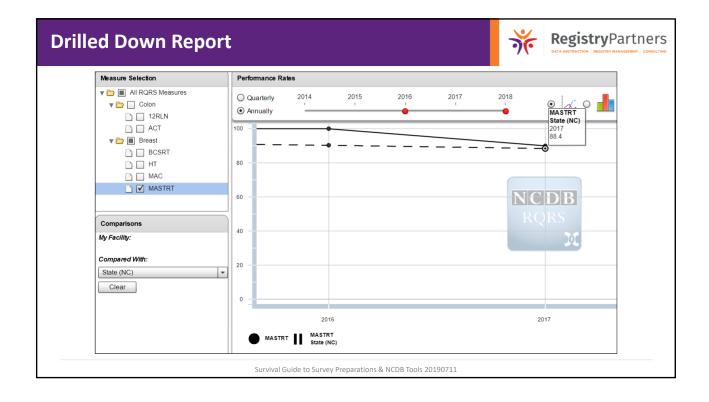


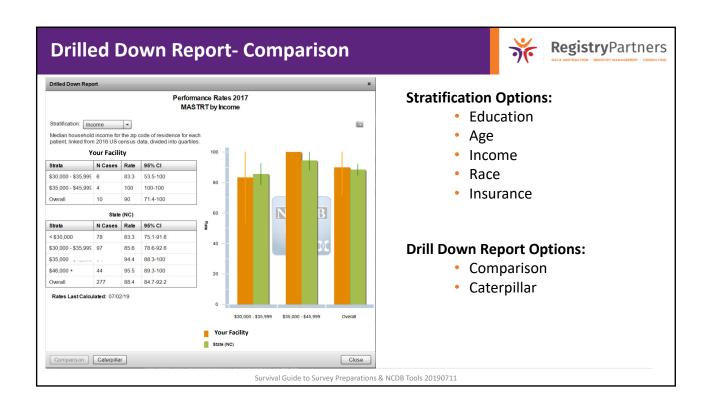


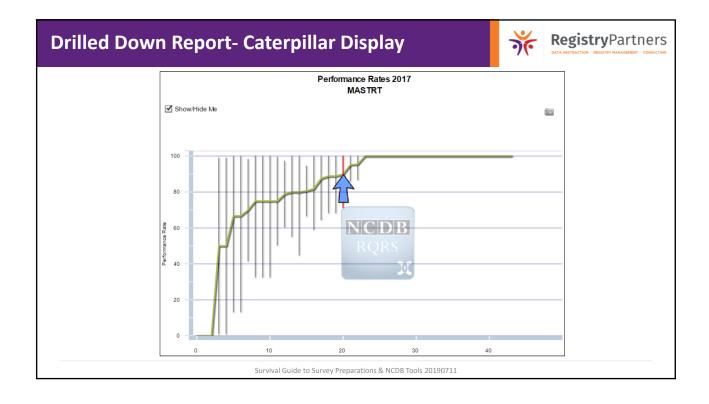


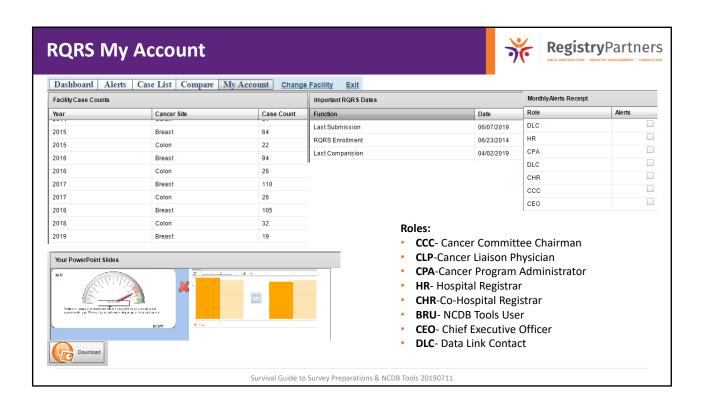




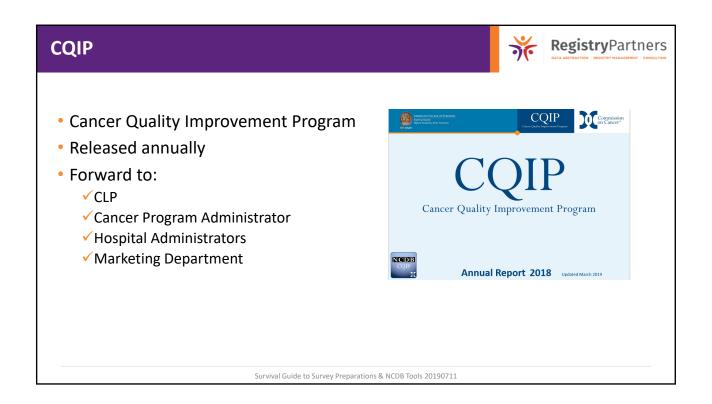








National Cancer DataBase (NCDB) Reporting Tools NCDB: Hospital Comparison Benchmark Reports NCDB: Survival Reports (V2) NCDB: Cancer Program Practice Profile Reports (CP3R) (v3) RQRS (v1.1) Cancer Quality Improvement Program (CQIP) Reports Annual Cancer Case Volume National Cancer DataBase (NCDB) Data Transmission NCDB: Submit/Resubmit Data Accepting v16 and v18 NCDB: Data Submission History and Edits NCDB: Authorize Third Party Submission NCDB Completeness and Default Overuse Reports



CQIP Includes...



- General Commission on Cancer information
- CoC Recommendations on the use of the **CQIP** Report
- Quality Measures for various cancer sites
- Mortality rates
- Major surgical resection data for selected
- Stage distribution for select cancer sites
- Distance traveled
- Treatment comparisons
- · Cancer program volumes
- In-migration and out-migration rates
- Insurance status data
- Survival data

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Slide #11: Quality Measure Reports

Slide #46: Surgical Volume and Unadjusted 30, 90 Day Mortality After Complex Operations

Slide #56: Survival Reports

Slide #71: Breast Cancer - Additional Reports

Slide #81: Colon Cancer - Additional Reports

Slide #89: Non-Small-Cell Lung Cancer (NSCLC) - Additional Reports

Slide #98: Prostate Cancer - Additional Reports

Slide #108: Melanoma of the Skin Cancer - Additional Reports

Slide #116: Commission on Cancer

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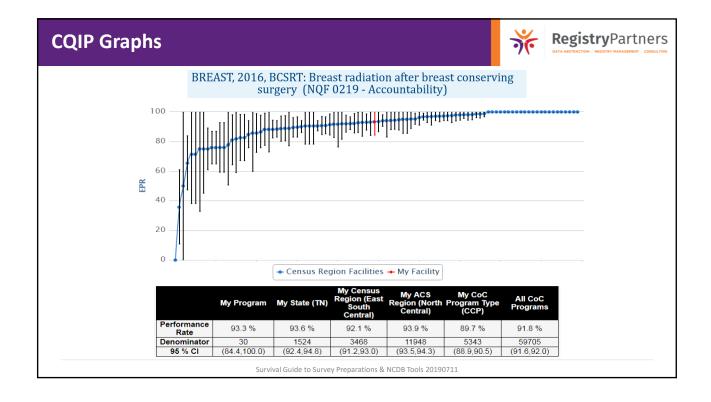
CQIP Recommended Uses

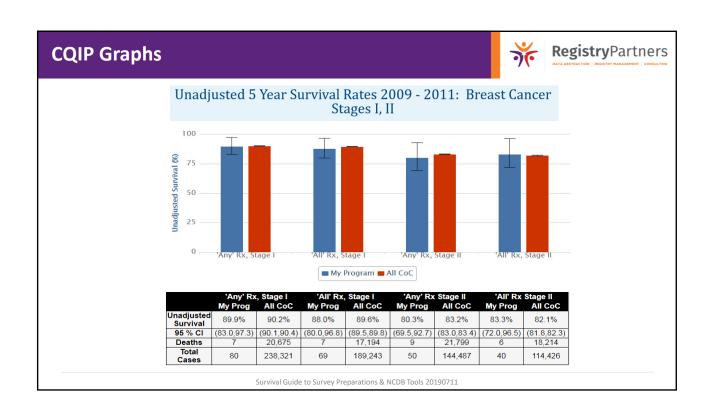


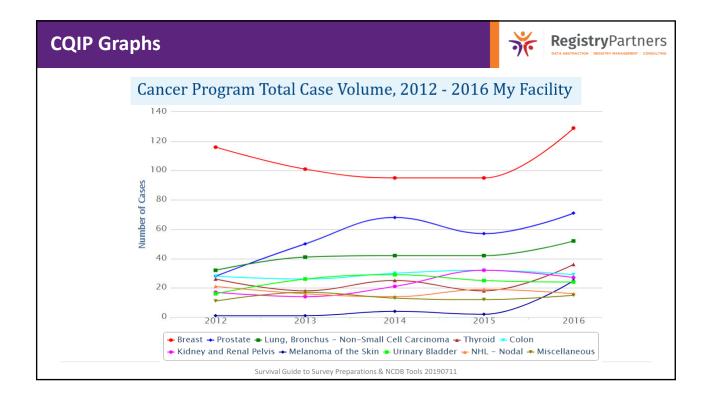


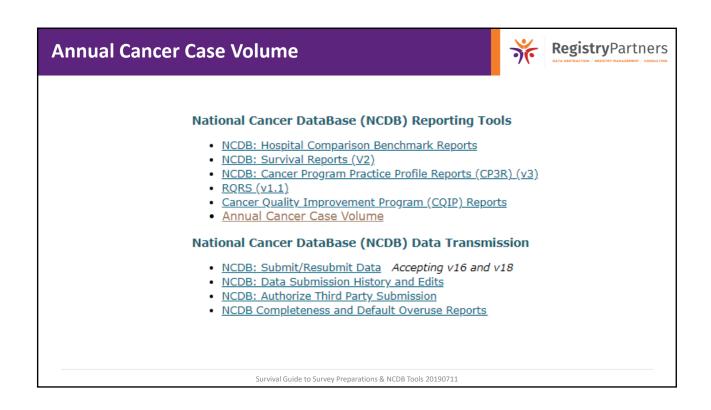


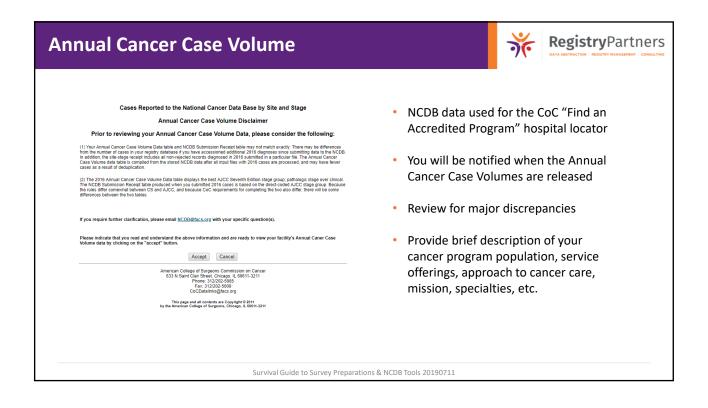
- Present and discuss at the Cancer Committee meeting
 - Major findings relevant to the cancer program
 - Recommended interventions for improvement of quality of cancer care
- Cancer Committee Leadership should present the report, major findings and recommendations to hospital leadership

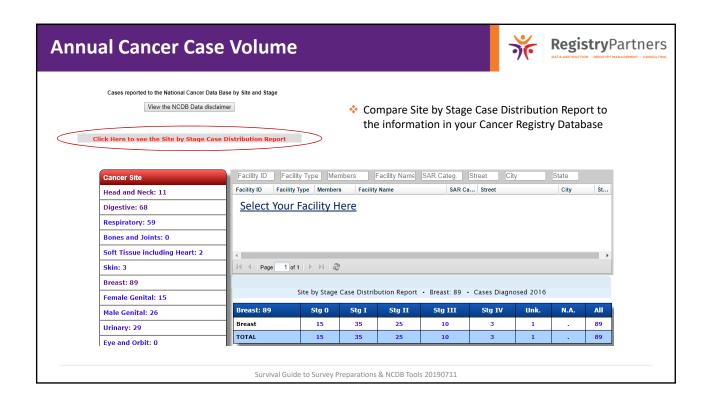


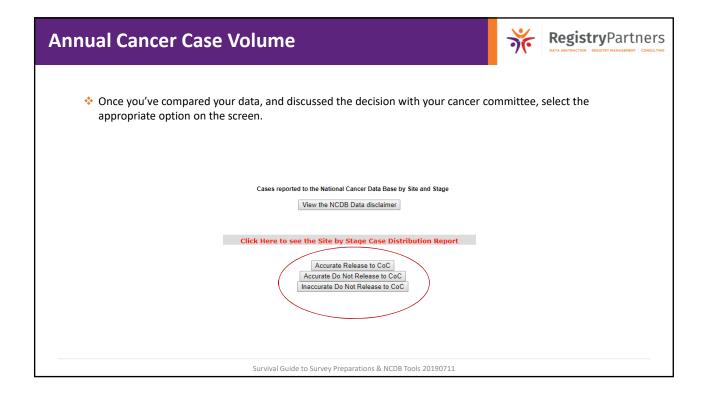


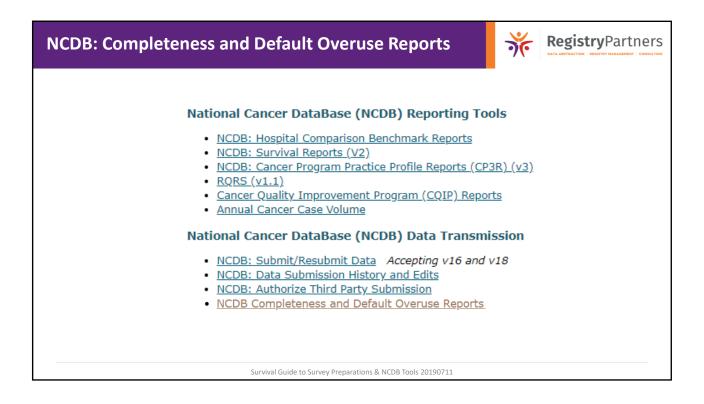


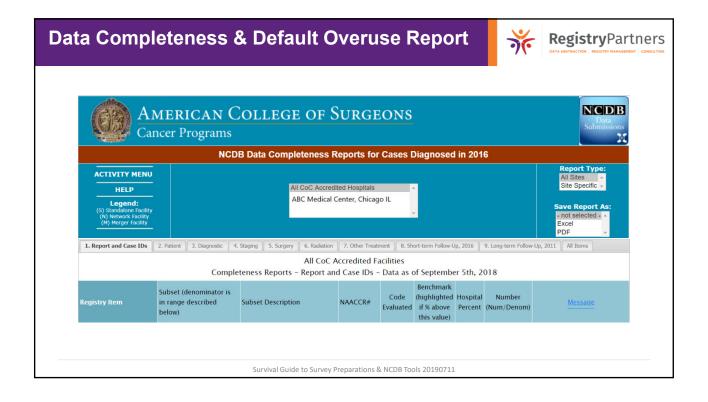


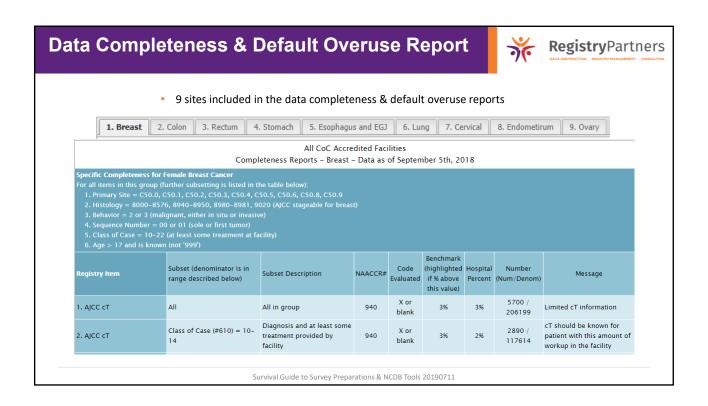


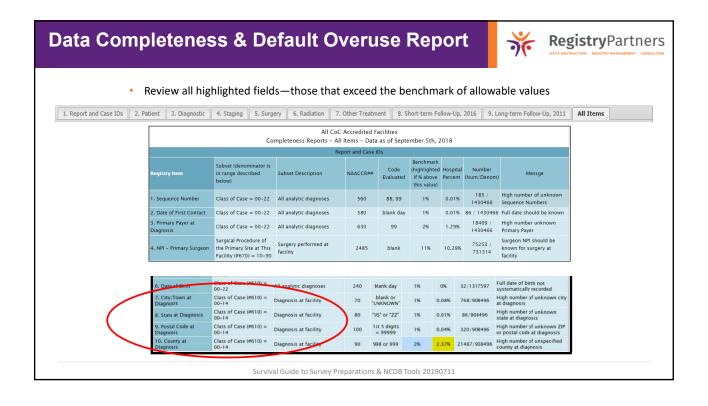


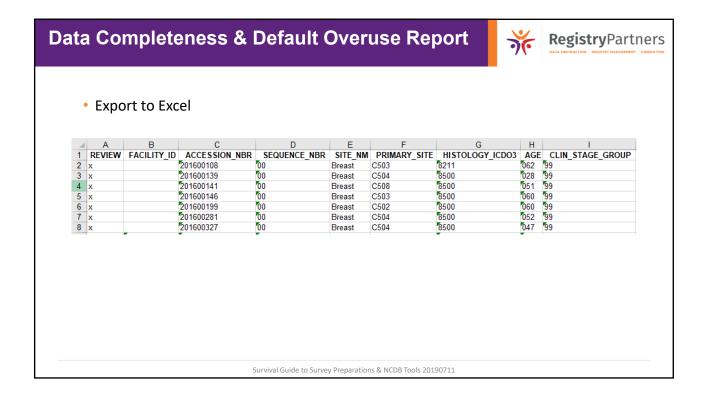












Public Reporting of NCDB Data RegistryPartners Request type **Tool/Source Approved Uses Reporting of NCDB Data** Reporting your program's benchmark information Benchmark Reports, CQIP Comparison of benchmark data to similar types of cancer programs **Benchmark Reports** Comparisons to cancer program's state, ACS region or all CoC Benchmark Reports, CQIP Annual or quarterly program compliance with Accountability and Quality Improvement, CP3R, RQRS, CQIP including measure comparisons. **Non-Approved Reporting of NCDB Data** Survival rates Survival Reports, CQIP Surveillance measures rates CP3R, RQRS, CQIP **RQRS** Year-to-date compliance rates CQIP Heat map Completeness of registry data Completeness and Default Overuse Report Survival Guide to Survey Preparations & NCDB Tools 20190711

Conclusion



- Use strategic planning when preparing a cancer program for survey
- Find the tools that work for you and your committees
- Hold all committee members accountable
- Support your Cancer Liaison Physician through education
- Know how to utilize all NCBD Reporting tools









- American College of Surgeons Cancer Programs: https://www.facs.org/quality-programs/cancer
 - Cancer Liaison Physician: https://www.facs.org/quality-programs/cancer/clp
 - Commission on Cancer: https://www.facs.org/quality-programs/cancer/coc
 - National Accreditation Program for Breast Centers: https://www.facs.org/quality-programs/napbc
 - National Accreditation Program for Rectal Cancer: https://www.facs.org/quality-programs/cancer/naprc
 - National Cancer Database: https://www.facs.org/quality-programs/cancer/ncdb
 - CoC Datalinks: https://web5.facs.org/Cancer/Account (login required)

Thank You For Your Attention!



RegistryPartners



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Coming UP...

Collecting Cancer Data: Solid Tumor Rules

• 08/01/2019

Collecting Cancer Data: Coding Pitfalls (last webinar of the season)

- 09/05/2019
- Guest Host: Janet Vogel, CTR

NAACCR

2019-2020 Webinar Series

10/3/19** Breast Wilson Apollo, CTR, Radiation Therapist and Jim Hofferkamp, CTR

11/7/19** Bladder Iris Chilton, CTR, Alberta Cancer Registry and Jim Hofferkamp, CTR

12/5/19 Base of Tongue/Head and Neck Wilson Apollo, CTR, Radiation Therapist and Jim Hofferkamp, CTR

1/9/20 **Prostate** Bobbi Matt, BS, RHIT, CTR State Health Registry of Iowa and Jim Hofferkamp, CTR

2/6/20* SSDI's: An In-Depth Look Jennifer Ruhl, CTR, Chair SSDI WG, Public Health Analyst NIH/NCI SEER, and Jim Hofferkamp, CTR

3/5/20** Abstracting and Coding Boot Camp Jim Hofferkamp, CTR

4/2/20 Melanoma Denise Harrison, CTR and Louanne Currence, CTR

5/7/20** Central Nervous System Denise Harrison, CTR and Louanne Currence, CTR

6/11/20* Esophagus Tonya Brandenburg, CTR Kentucky Cancer Registry

7/9/20** Navigating the 2020 Survey Application Record (SAR) Cynthia Boudreaux, LPN, CTR, Owner/Consultant CB Professional Abstracting

8/6/20* Corpus Uteri Denise Harrison, CTR and Louanne Currence, CTR

9/3/20** Coding Pitfalls Janet Vogel, Compliance and Quality Auditor/Educator-Cancer Registry himagine solutions, inc.

https://www.naaccr.org/cancer-registry-surveillance-webinar-series/#purchase

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CE Certificate Quiz/Survey

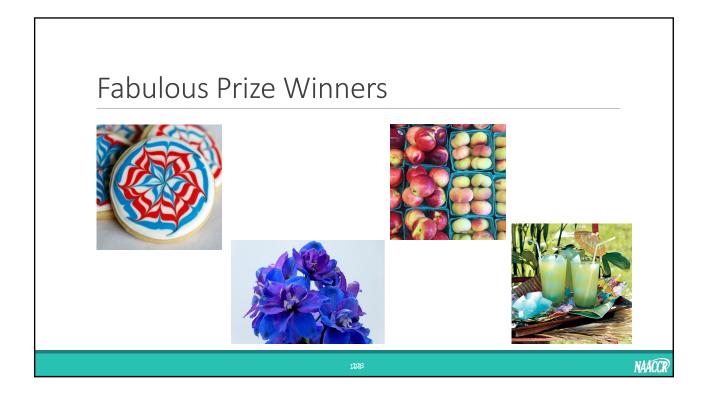
Phrase

Benchmark

Link

https://www.surveygizmo.com/s3/5104850/A-Registrars-Defense-to-ACoS-Accreditations

NAACCR



Thank you!

SARA MOREL <u>SARA.MOREL@MIDMICHIGAN.ORG</u>

COURTNEY JAGNEAUX <u>COURTNEYJAGNEAUX@REGISTRYPARTNERS.COM</u>

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