Ready, Set, Go ... Call for Data & v22 Changes







Presenter



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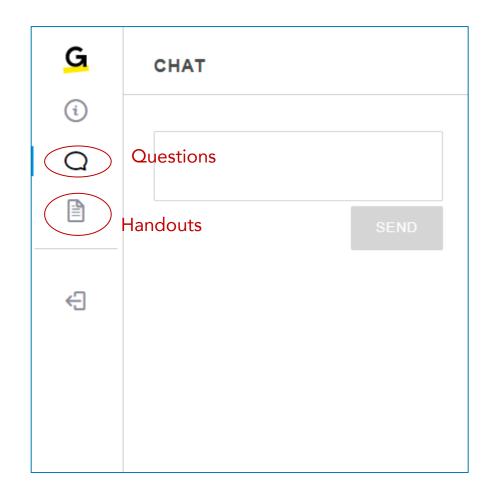
Housekeeping

- If you have trouble hearing the presentation and are listening through your computer, please check and adjust your volume settings. There will not be a call in feature for this webinar. If your speakers are having issues you may be able to use headphones with your computer.
- The webinar will be approximately 1 hour in length
- CE certificates will be sent to attendees later this week to the email address you provided during registration for this webinar
- A recording of this webinar, along with handouts and the Q&A will be made available to attendees in the coming days



Navigating GTW

- Download handouts and ask questions from the GTW control panel
- Ask questions! All questions will be answered in a follow up Q&A document





Objectives

- Explore resources available for implementation of 2022 changes
- Familiarize participants with new and changed data items from NAACCR and STORE
- To familiarize CRStar users with the processes involved in answering to the 2022 NCDB Call for Data
- To offer suggestions for Best Practices to help prepare for the Call for Data
- To demonstrate the use of the CRStar exporting application and the GenEDITS Plus 5 software











CoC 2022 Call for Data





NCDB Call for Data

• Information can be found here: https://www.facs.org/quality-programs/cancer/ncdb/call-for-data

NCDB Call for Data

Registrars

Information for Registrars

Data Submission Information

How to Submit Data

GenEDITS Plus 5—Download and install from the CDC website

CoC Program Standard 6.4

CoC Program Standard 6.4 Compliance

Schedule for NCDB Data Submission

Tool-NCDB Call for Data Submission Count Tracking

Configuration/Setup

Metafile and CoC User Dictionary

NCDB and RCRS v18 Metafile

NCDB and RCRS v21 Metafile

NCDB and RCRS v22 Metafile

V21 CoC User Specific Dictionary for NPI Physician Confidential Data

V22 CoC User Specific Dictionary with NPI and Breast Surgical Codes

Layouts



Data Submission Information

- Submission Window opens on 3/1/2022 and closes on 3/31/2022 at 11:59 CST
- Required Submission Existing Programs
 - Analytic cases
 - All <u>new</u> cases diagnosed on or after 2006 through 2020
 - All updated/changed cases diagnosed on or after 2006 through 2019 (Those previously submitted in prior call for data exports)
- Required Submission New Program
 - All analytic cases diagnosed on or after 2006 or their first accredited date (whichever is shorter) through 2020



Prep Work

- Review instructions on NCDB website
- Review for complete staging, treatment and follow-up information
- Review FIN to verify valid number
- Carefully choose your file name so that you can easily find file for submission

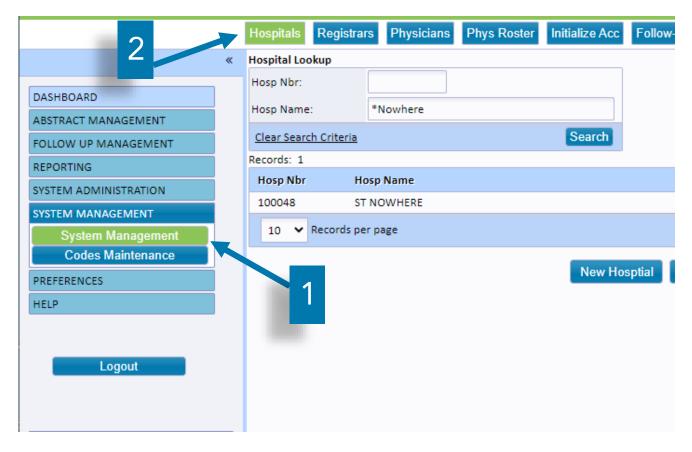
 NCDB Call for Data
- Zip files will NOT be accepted



11

Registrars

Prep Work- ACoS FIN number verification



Hospital Name:	ST NOWHERE	
Contact:		
Phone Nbr:	(999) 555-1123	
Rpt Hosp Id:		
St Name Add:	123 MAIN ST	
Acos Hosp Id:	12345678	
NPI:	1699722728	
Region:		
	1699722728	



Prep Work: FIN and NPI numbers

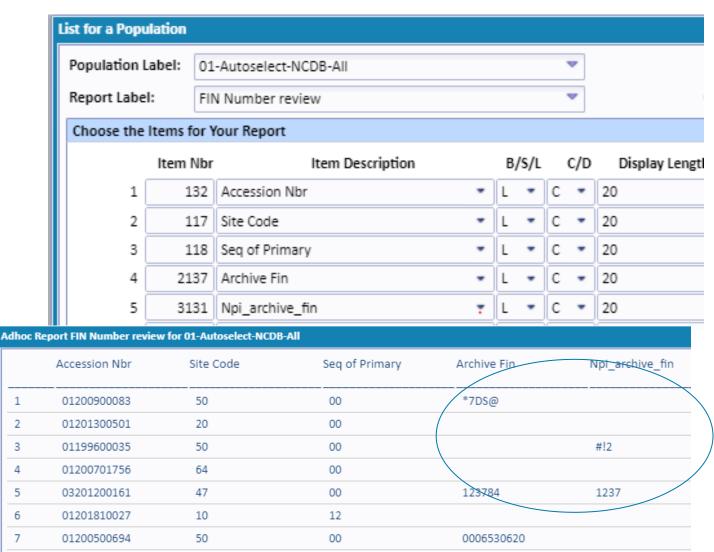
2

3

4

- This task can be done after selecting your export population
- Use List for a Population under Adhoc (do not reselect the auto-selected population
- List the fields to review including **Archive FIN and NPI Archive FIN**

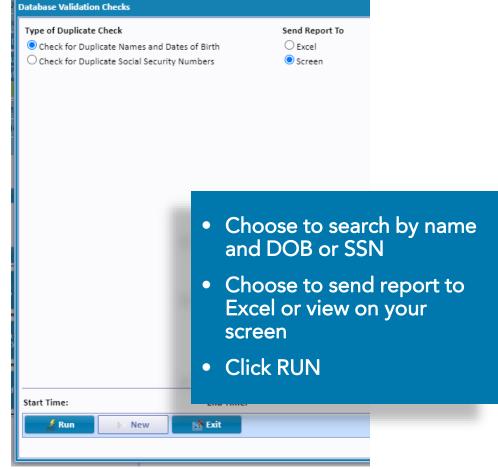




Best Practices: Database Validation

Perform Database Validation Check/Eliminate Duplicates

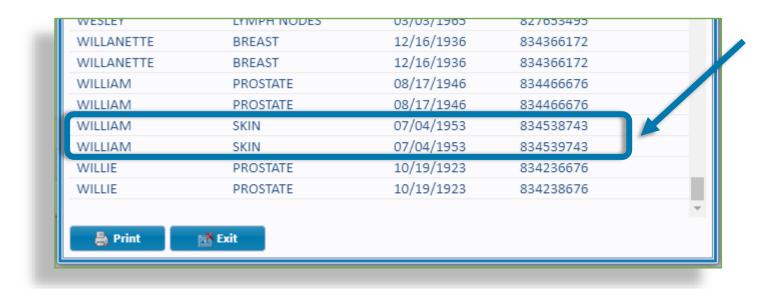






Best Practices: Database Validation (con't)

 Identify any duplicates. Investigate them in hospital EMR to determine which data is accurate, then combine the patients.

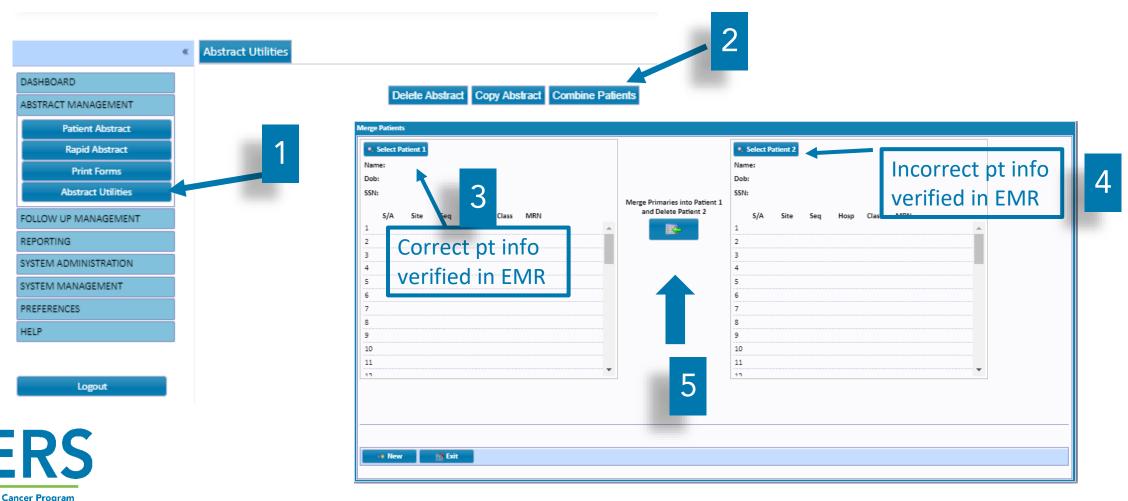




Best Practices: Database Validation (con't)

Combine patients

Management & Analytics



Best Practices: Follow Up

- Before starting the Call for Data process, get ahead by doing a thorough job of Follow-Up
 - Include applicable follow-up, recurrence, death and updated treatment data
 - Accurately input
 - Edits run
- Serves three purposes:
 - Ensures the most updated cases are submitted
 - Less errors in Follow-Up data
- No harm in pausing FU activities when it's time to submit



Best Practices: GenEdits Plus 5 & Metafile

- The metafile for the 2020 Call for Data is: NCDB v21b
- It is important that everyone on your team has the correct metafile before trying to reconcile any errors identified by the GenEDITS Plus 5 software
- Utilize one person to run the GenEDITS
 - More than one person can process the edits
- Assign one person to run final GenEDITS and submit data



Best Practices: General

- File storage
- Pause all Follow-up and Import activities while completing the final run of edits and submission
- RCRS Platform
 - Make sure the appropriate people have access
 - Check your facility profile for accurate information





Common Errors

- Conflicts in Follow-Up info
 - Recurrence type 70 and Cancer status 1
 - Recurrence type blank and Cancer status 2
 - Recurrence date entered with no Recurrence type
- Conflicts in Class of Case info
 - Class 00 without Referred to facility
 - Class 20-22 without Referred from facility
- Conflicts in Staging data
 - Having both (p) values and (yp) values on same case



What could cause my file to be rejected?

- Your FIN is incorrect in the transmission file or more than one FIN in the file
- File is not formatted for v21 or v22
- Submitted incorrect state export file instead of NCDB export
- More than one period in the file before extension (.xml)
- Special characters in file name. Only hyphen (-) and underscore
 (_) are accepted



GenEDITS Plus 5

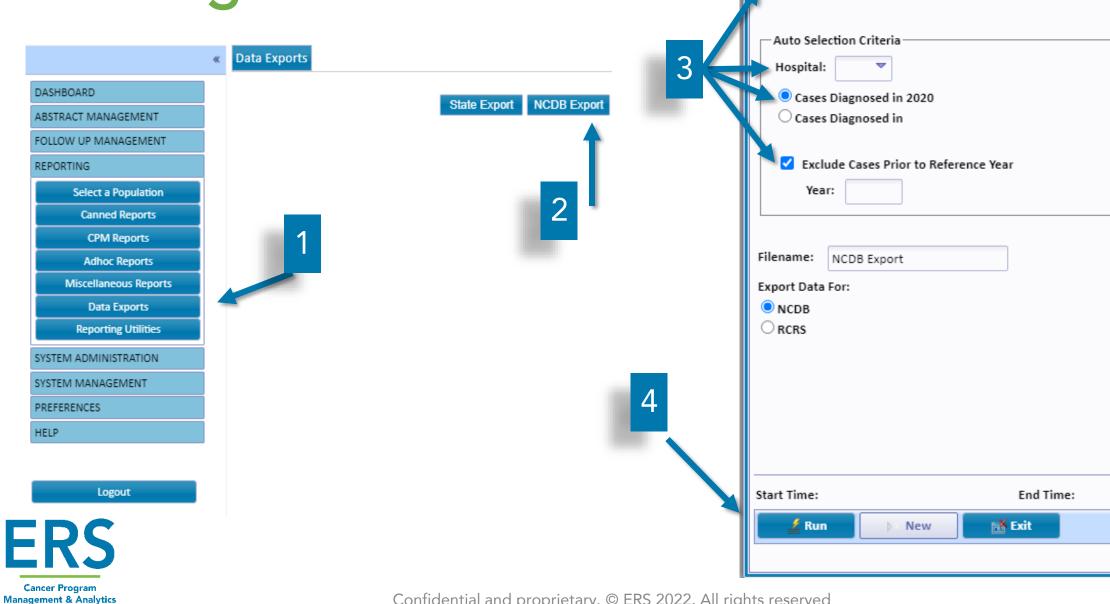
Instructions
 https://www.facs.org/-/media/files/quality-programs/cancer/ncdb/2021_genedits_plus_5_installation_v18_and_v21.ashx

- Download <u>https://www.cdc.gov/cancer/npcr/tools/edits/edits50.htm</u>
- Metafile Installation
 https://www.facs.org/quality-programs/cancer/ncdb/call-for-data

Hints: Hospital Firewalls can interfere with download.



Creating Your Data File

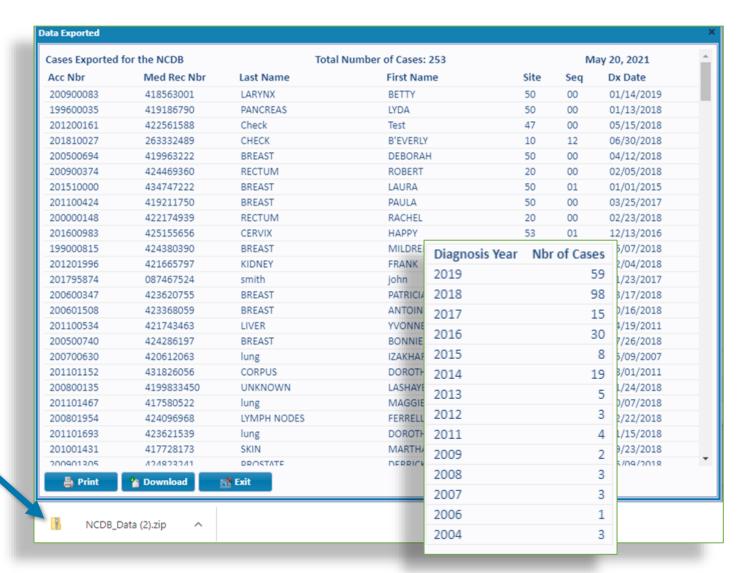


Export NCDB Data

Auto Select Use Existing Population

Creating Your Data File

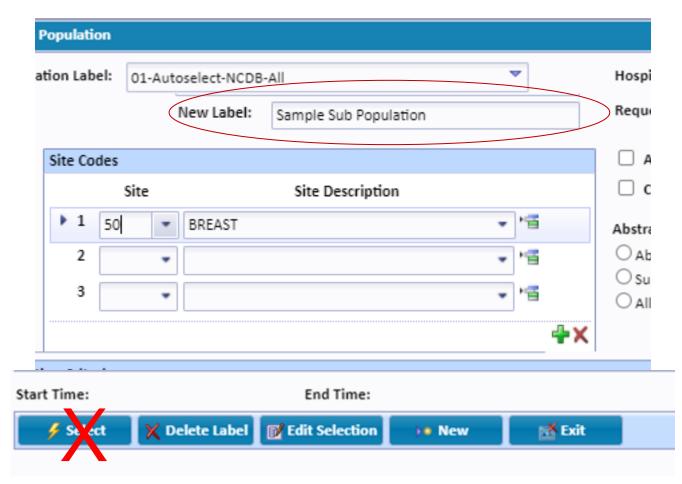
- The 2020 Export file is created and a list of cases is displayed
- At the bottom of the list, a summary of the contents is displayed
- The export list can be printed or saved (printed) as a PDF for reference
- Click on the download indicator at the bottom left of your screen to go to the file





Your Auto-Selected Population

- If you need to run other reports on the auto-selected population, it is available in your Population Label drop-down menu, BUT, if you open it, there will be no selection criteria listed
- Do NOT reselect the population here, it will come up with zero cases
- You can, however, create a new label and run a subpopulation as shown in this example:

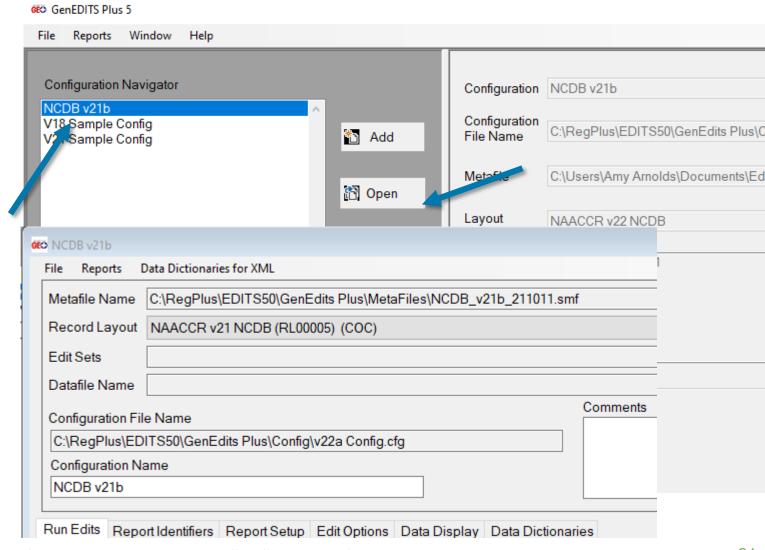






Cleaning Up Your Data

- Open GenEDITS Plus 5
- v21b Configuration
- Click Open

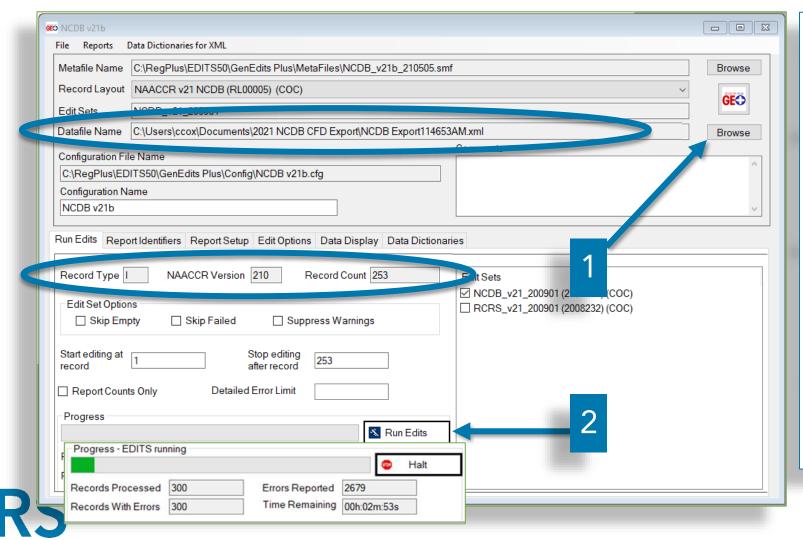




Cleaning Up Your Data

Cancer Program

Management & Analytics



- 1. Click Browse & choose your data file
- Confirm that it populates in the Datafile Name box
- Confirm that your Record type, NAACCR version and Record Count are correct
 - 1. Run Edits button should now be activated

Cleaning Up Your Data (con't)

- GenEDITS Plus 5 generates your edit reports as you specified
- The summary report shows you how many of each edit type you have
- The detail report shows you each case that has edits with identifying information so you can clean those edits up in CRStar



Cleaning Up Your Data

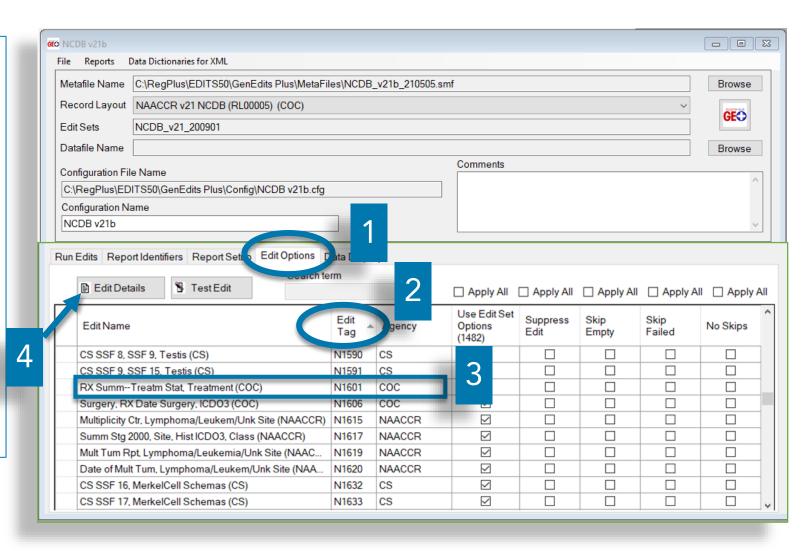
 Edit numbers can be looked up in GenEDITS Plus 5 to help clarify what all of this means ...





Cleaning Up Your Data - Finding 4 digit Edit

- Go to Edit Options tab
- 2. Double-click on Edit tag to sort the edit tag numbers numerically
- 3. Find the edit in the list and highlight it
- 4. Click Edit Details





30

Cleaning Up Your Data

Look for keywords like "must equal", "cannot be", etc



Edit Name: RX Summ--Treatm Stat, Treatment (COC) (N1601)

Agency: COC

Description Admin Notes Edit Logic

The purpose of this edit is to verify that RX Summ--Treatment Status and treatment fields are coded consistently.

This edit is skipped under the following conditions:

- a. RX Summ--Treatment Status is blank
- b. Date of Diagnosis is blank (unknown) or invalid
- If any of the treatment fields indicate treatment given, then RX Summ--Treatment Status must equal 1 (treatment given). Treatment is considered "given" if any of the following is true:

RX Summ--Surg Prim Site = 10-90

RX Summ--Surg Oth Reg/Dis = 1-5

RX Summ--BRM = 01

RX Summ--Chemo = 01-03

RX Summ--Hormone = 01

RX Summ--Transplnt/Endocr = 10-40

RX Summ--Other = 1, 2, 3, 6

Reason for No Radiation = 0

2. If Treatment Status = 1 (treatment given), at least one of the following fields must equal treatment given. Treatment is considered "given" if any of the following is true:

RX Summ--Surg Prim Site = 10-90

RX Summ--Scope Reg LN Sur = 1-7

RX Summ--Surg Oth Reg/Dis = 1-5

RX Summ--BRM = 01

RX Summ--Chemo = 01-03

RX Summ--Hormone = 01

RX Summ--Transplnt/Endocr = 10-40

RX Summ--Other = 1, 2, 3, 6

Reason for No Radiation = 0

Cleaning Up Your Data

- Run edits on every case you touch during the cleanup process
- Review and clear all edits
- Run export again and pass through GenEDITS
 - · If edits are clear, data can be submitted
 - If edits aren't clear they will need to be worked on then re-exported until clear



Submitting the Call for Data

- Instructions to Submit Data: https://www.facs.org/-
 /media/files/quality-programs/cancer/ncdb/how-to-submit-data.ashx
- Questions or Issues? ACSTechsupport@iqvia.com



Determining Submission Completeness

- Keep track of how many cases were in each Export batch you created
- Errors & Rejected Cases Any rejected cases are considered not submitted. Correct them and re-submit prior to the deadline 3/31/2021
- Compare to the number of cases received by the NCDB after submitting



Determining Submission Completeness

The NCDB uses this formula and these percentages to determine if a facility's data submission meets the expectations for submission based on year.

The NCDB monitors the frequency of updated cases that were submitted by facilities for the earlier years, and evaluates the completeness of reporting of cases for these diagnosis years based on a thorough analysis of data submitted from all facilities to determine an expected number of cases:

Diagnosis Years 2004-2018

The expected number of cases for each facility is based on a percentage that is calculated as follows:

cases submitted for the diagnosis year (e.g., 2017)

cases submitted for the most recent diagnosis year (e.g., 2018)

X 100 = ZZ%

Percent of case submission expectations for Standard 5.5 Compliance:

Year	Percentage
2018	100
2017	85
2016	50
2015	30
2014	25
2013	20
2012	15
2011	10
2004-2010	5



Determining Submission Completeness

NCDB Call for Data Submission Count Tracking by Diagnosis Year

Call for Data 2022					
Dx year	CFD 2022	CFD 2021	% Annual		
	# Cases	# Cases	Caseload		
First dx year	Enter # of	Enter # of			
	cases	cases	#VALUE!		
	submitted for	submitted for			

Enter the number of cases submitted for each diagnosis year for your program's annual caseload will be automatically calculated

Call for Data 2022					
Dx year	CFD 2022	CFD 2021	% Annual		
	# Cases	# Cases	Caseload		
2020					
2019			#DIV/0!		
2018			#DIV/0!		
2017			#DIV/0!		
2016			#DIV/0!		
2015			#DIV/0!		
2014			#DIV/0!		
2013			#DIV/0!		
2012			#DIV/0!		
2011			#DIV/0!		
2010			#DIV/0!		
2009			#DIV/0!		
2008			#DIV/0!		
2007			#DIV/0!		
2006			#DIV/0!		



Responsibilities During the Call For Data (Vendor Responsibilities)

- Support tickets will be answered in the order received. We may need to limit the amount of time spent on each call
- ERS will not provide support for GenEDITS Plus 5 software
- ERS will gladly assist with any *software related edits*. However, we are unable to provide clean-up services for data entry errors
- ERS will assist with global data changes if applicable and the amount of any one edit is over 75 errors.
 - The edits below require manual review and globals are not indicated:
 - Recurrence Type 1st, Cancer Status (COC)
 - Date Case Completed CoC, Date of Diagnosis (NCDB)
 - Rx Summ Treat Stat, Date 1st Crs RX COC (COC)



Responsibilities During the Call for Data (Client Responsibilities)

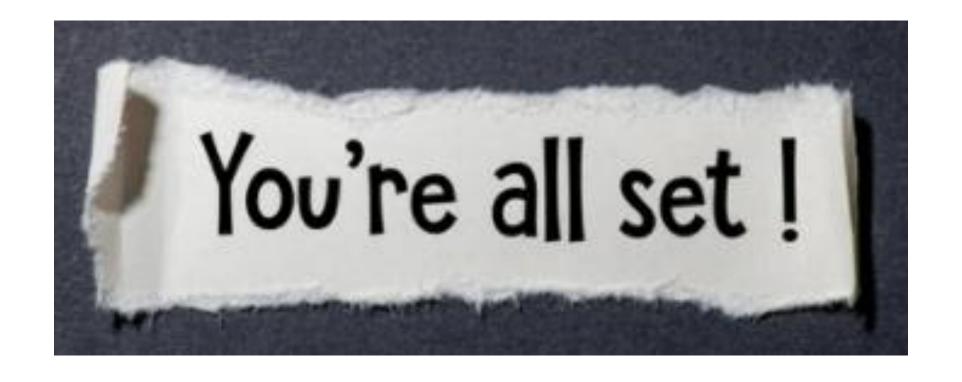
- Please submit a support ticket if assistance is needed with an ERS issue.
 Please do not email or call the support staff directly
- For assistance with GenEDITS Plus 5 software, contact the CDC as this software is developed and supported by the CDC. For assistance with the NCDB submission portal, contact the NCDB
- During this time, it is imperative for Registrars and Follow-Up clerks to run NCDB edits after entering Follow-Up and subsequent treatment information. If new FU is entered, the case becomes eligible for this Call for Data. Make sure it passes edits
- · A backup of the database must be done prior to any global changes



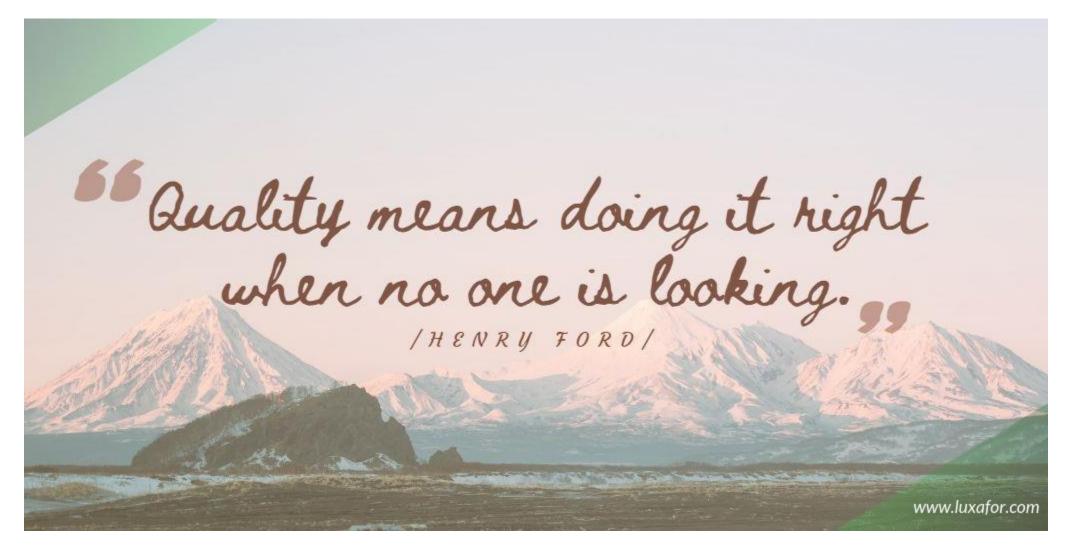
Quick Review

- Get organized. Review the 2022 Call for Data Requirements
- Do some preliminary data reviews
- Check for incorrect FIN and NPI numbers
- Install GenEDITS Plus 5
- Download the NCDB v21b metafiles and import them into GenEDITS
- Create the NCDB 2022 CFD Configuration
- Create the 2020 and earlier Data Export files in CRStar
- Run edits on the Export files in GenEDITS Plus 5 (repeat until clean) and save a copy of the final clean report Just Breathe
- Submit data through NCDB Datalinks portal
- **Determine Submission Completeness**











Resources - Updated Quick Links

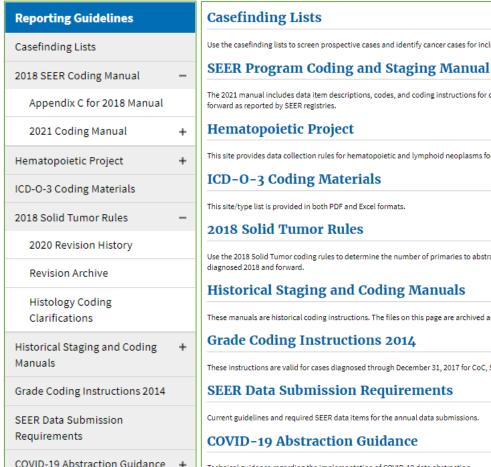
- Links will go to the most current version of the standard setters' sites and manuals
- Use caution to make sure you are looking at the correct version for the year of the case being abstracted







SEER Information Hub





NATIONAL CANCER INSTITUTE

Surveillance, Epidemiology, and End Results Program

Use the casefinding lists to screen prospective cases and identify cancer cases for inclusion in the registry,

The 2021 manual includes data item descriptions, codes, and coding instructions for cases diagnosed January 1, 2021 and forward as reported by SEER registries.

Hematopoietic Project

This site provides data collection rules for hematopoietic and lymphoid neoplasms for 2010+

ICD-0-3 Coding Materials

This site/type list is provided in both PDF and Excel formats.

2018 Solid Tumor Rules

Use the 2018 Solid Tumor coding rules to determine the number of primaries to abstract and the histology to code for cases diagnosed 2018 and forward.

Historical Staging and Coding Manuals

These manuals are historical coding instructions. The files on this page are archived and provided for reference purposes only.

Grade Coding Instructions 2014

These instructions are valid for cases diagnosed through December 31, 2017 for CoC, SEER, and NPCR

SEER Data Submission Requirements

Current guidelines and required SEER data items for the annual data submissions.

COVID-19 Abstraction Guidance

Technical guidance regarding the implementation of COVID-19 data abstraction.

- https://seer.cancer.gov/registr ars/quidelines.html
- Each section has information / links for previous versions and 2022 version



v22 Implementation Updates

NAACCR 2022 Implementation Guidelines & Recommendations

https://www.naaccr.org/wp-content/uploads/2021/12/2022-Implementation-

Guidelines_20211129-1.pdf

Reminder: Most v22 changes discussed apply to cases dx 1/1/2022 and later



v22 - New Data Items

- Tobacco Smoking Status Patient Screen
- SSDI

Item #	SSDI Name	Schema
3955	Derived Rai Stage*	Lymphoma CLL/SLL
3956	p16**	Cervix V9
3957	LN Status Pelvic***	Cervix 8th, Cervix V9, Vagina, Vulva
3958	LN Status Para- Aortic***	Cervix 8th, Cervix V9, Vagina
3959	LN Status Femoral- Inguinal***	Vagina, Vulva



v22 - New Data Items - CoC

- CoC -Macroscopic
 Evaluation of Mesorectum
- CoC Breast Surgical (2022 only)
 - RX Hosp Surg Breast
 - RX Summ Surg Breast
 - RX Hosp Recon Breast
 - RX Summ Recon Breast

Surgical Screen

https://www.facs.org/qualityprograms/cancer/events/ondemand/canswerforum-live



v22 - Changed Data Items

- Phrases -
 - Examples: API to DLL; Alaska Native, etc.
- SSDI Cases Dx 1/1/2018 forward
 - Blanks
 - New Codes
 - Schemas: Esophagus, Esophagus Squamous, Stomach, Prostate, Kidney Parenchyma, Pancreas, Plasma Cell Myeloma, Lymphoma CLL/SLL, Plasma Cell Myeloma
 - Removed
 - Schemas: Cervix 8th, Cervix V9, Vulva, Vagina



v22 - Solid Tumor Rules



- The 8 site groups have been updated:
 - New histologies, codes & terms
 - Updated equal/equivalent and not equal/equivalent terms
- New section added "Changes from 2018 Solid Tumor Rules" for Colon and Head & Neck
- Timing requirement updates for Colon
- Solid Tumor General instructions apply to all sites



v22 - Solid Tumor Rules (con't)



- Head & Neck
 - p16 can be used to determine HPV positivity for squamous cell carcinoma
 - 1/1/2022 and forward Non-keratinizing squamous cell ca, HPV pos is 8085 for sites listed in Table 5 ONLY; non-keratinizing squamous cell ca, NOS = 8072
 - 1/1/2022 and forward Keratinizing squamous cell ca, HPV neg = 8086 for sites listed in Table 5 ONLY; keratinizing squamous cell ca, NOS =8071





v22 - ICD-O-3

- Review v22 Implementation Guide for updates
- Reportable
 - Clear cell papillary renal cell carcinoma has been reclassified as /1 but remains reportable for Urinary system & Male Genital organs
 - Low-grade appendiceal mucinous neoplasm (LAMN) now /2 or /3 reportable
- Not Reportable
 - High grade dysplasia /2 (colon)
 - Adenoca in situ, HPV-associated /2 (cervix)
 - Adenoca in situ, HPV-independent /2 (cervix)



v22 - Extent of Disease (EOD)



- Cervix Sarcoma new schema
- Soft Tissue Other is now split
 - Soft Tissue Rare
 - Soft Tissue Other
- Pleural Mesothelioma
 - 05 is a new code for positive pleural effusion only



v22 - Metafile



 v22 metafile will only be available for selection if your state is accepting v22 NAACCR layout for submissions

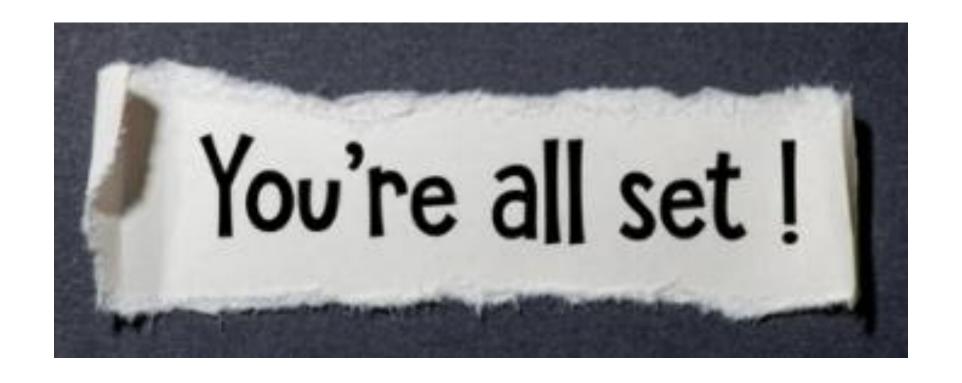


v22 - CoC



- STORE v2022 for all cases dx on/after 1/1/2022
- COVID-19 Fields not required for cases dx on/after 1/1/2022
- Histologies no longer collected
 - 8210/2 Adenomatous polyp, high grade dysplasia (C160 C166, C168-C169, C170-C173, C178-C179)
 - 8211/2 Tubular adenoma, high grade
 - 8261/2 Villous adenoma, high grade
 - 8263/2 Tubulovillous adenoma, high grade
 - 8483/2 Adenocarcinoma in situ, HPV-associated (C530-C531, C538-C539)
 - 8484/2 Adenocarcinoma in situ, HPV-independent, NOS C530-C531, C538-C539)
 - 8590/1 Uterine tumor resembling ovarian sex cord tumor
 - 9200/1 Osteoblastoma
 - 9261/1 Osteofibrous dysplasia-like adamantinoma







Converted Data Items

- NAACCR v22 upgrade required conversion of several previously coded fields
 - Some require manual review
 - Specifications are in the 2022
 Implementation guidelines
 Appendix B
 - ERS has provided you with population lists of cases that had conversions executed

Schemas that require review:

- 13.1 AJCC [995] changes within Schema ID [3800] (Recalculation) Schema ID changed and cases are now eligible for AJCC staging. AJCC T, N, M and Stage Group should be reviewed and restaged
- 13.2 Schema ID [3800] = 00421 (Soft Tissue Abdomen & Thoracic) C474
 (Recalculation) Schema ID changed. AJCC T, N, M and Stage Group should be reviewed and restaged
- 13.3 Schema ID[3800] = 00450 (Soft Tissue Rare), 00459 (Soft Tissue Other) New Schema (Recalculation) Schema ID changed. AJCC T, N, M and Stage Group should be reviewed and restaged
- 13.4 Schema ID [3800] = 00450 (Soft Tissue Rare) C530-C539 (Recalculation)
 Schema ID changed. AJCC T, N, M and Stage Group should be reviewed and restaged
- 13.5 Schema ID [3800] = 00528 (Cervix Sarcoma) New Schema (Recalculation)
 Schema ID changed. AJCC T, N, M and Stage Group should be reviewed and restaged
- 13.10 Schema ID [3800] = 09520 (Cervix Version 9) and p16[3956] Text will need to be checked for p16 test results and SSDI updated accordingly



Managing the Conversion Cases

Populations have been created for you to use to manage the manual review process for cases that require it

Search: v22	Search All			
Hospital Code	Population Label 📤	Request Date	Requested By	Nbr Cases
	V22 Schema resets and manual review of newly eligible Histologies	01/27/2022	Automated v22	21
	v22 SchemalD 00421 Topo C474 Review	01/27/2022	Automated v22	21
	v22 SchemalD 00450 Topo C530-C539 Recalculation	01/27/2022	Automated v22	75
	v22 SchemalD 00528 New Schema Recalculation	01/27/2022	Automated v22	17
	v22 SchemalD 00700 Histo 9222_ to SchemalD 00450 or 00459	01/27/2022	Automated v22	35
	v22 SchemalD 09520 p16 Manual Review	01/27/2022	Automated v22	38

IMPORTANT: Use the populations in reports as they are.

DO NOT RESELECT THE POPULATIONS!

Reselecting will select 0 cases.



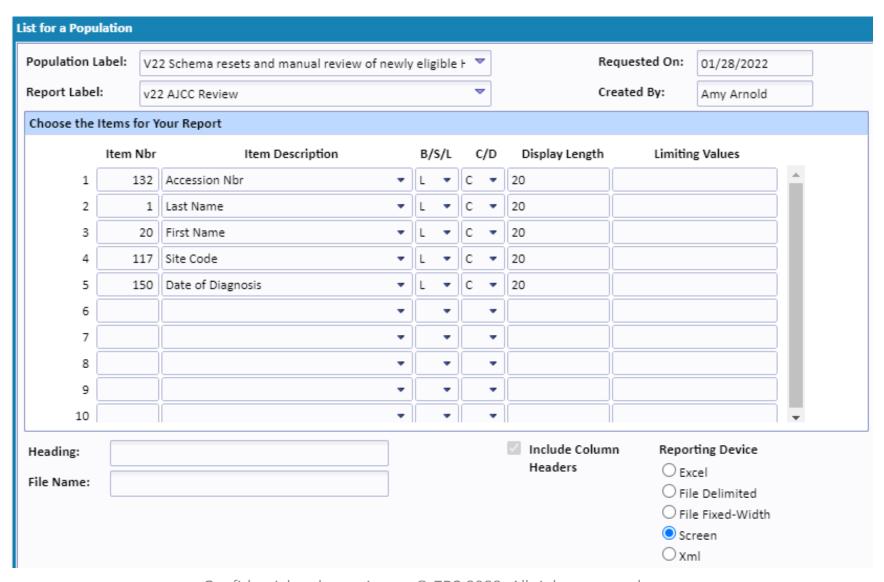
Managing the Conversion Cases

- Use the populations we created for you to generate Adhoc reports
- Review the cases in each report according to Appendix B in the NAACCR Implementation Guidelines

Section Number	Population Name	Items to Review
13.1a	V22 Schema resets and manual review of newly eligible Histologies	AJCC T,N,M & Staging Group all categories
13.2	V22 SchemalD 00421 Topo C474 Review	AJCC T,N,M & Staging Group all categories
13.3c	V22 SchemalD 00700 Histo 9222_ to SchemalD 00450 or 00459	AJCC T,N,M & Staging Group all categories
13.4	V22 SchemalD 00450 Topo C530- C539 Recalculation	AJCC T,N,M & Staging Group all categories, SSDI
13.5	V22 SchemaID 00528 New Schema Recalculation	AJCC T,N,M & Staging Group all categories , SSDI
13.10	V22 SchemalD 09520 p16 Manual Review	p16 SSDI, Lab/Path Text

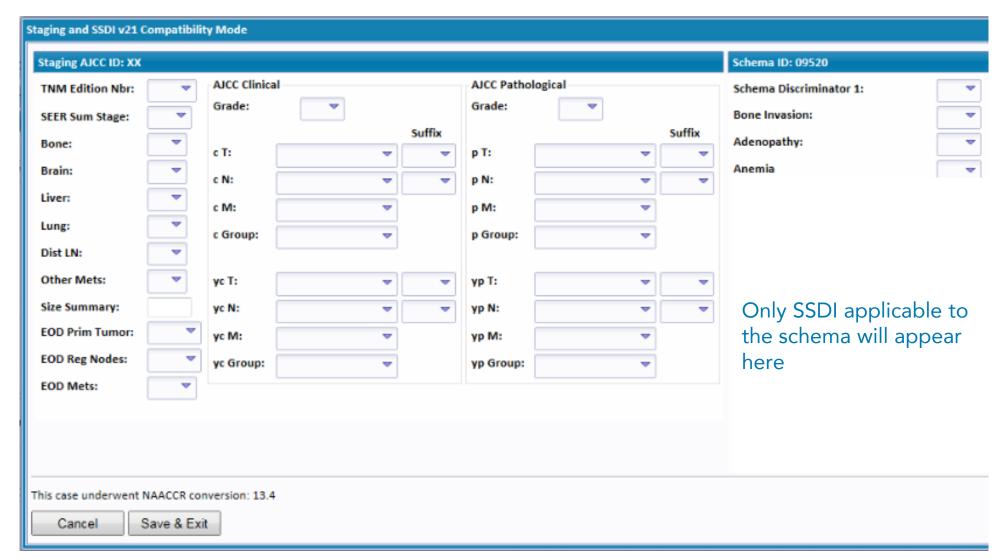


Suggested Report Review #1 - AJCC Review





v21 Compatibility Mode









Resources

- NAACCR 2022 Implementation Guidelines and Recommendations
 https://www.naaccr.org/wp-content/uploads/2021/12/2022-Implementation-Guidelines_20211129-1.pdf
- SEER Information Hub https://seer.cancer.gov/registrars/guidelines.html
- NCDB Call for Data https://www.facs.org/quality-programs/cancer/ncdb/callfor-data
- Contact Information
 - ncdb@facs.org questions regarding data submission
 - ACSTechsupport@iqvia.com RCRS login or upload issues





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