



# Quality Assessment of Cancer Registry Data

This presentation will be about the data – what registrars collect, code and record in the cancer registry database. Some of the tools that will be discussed are most often used to assess quality of patient care but we will be using these tools to help assess quality of data.

I am not an EXPERT – I am a registrar.





# Change

Cancer Registry is changing

Cancer Registry has changed multiple times before

Cancer Registry will continue to change

Processes and operations must change in order to meet the goals and outcomes of our work



## Innovation is not

Using the same Power Point Slide, 2X in a row!

Doing the same old thing the same old way

Always BIG

Always successful or instantaneous



SECTION 1

### Innovation is

Intentional yet creative

Calm or swirly

 $BIG \ or \ MEDIUM \ or \ small \ or \ even \ {}_{minuscule}$ 

Successful and yet may still need massaging

Cooperative . . . Go with the flow

Scary and exciting



#### SECTION 1

Initially ER and PR were recorded only in text as registrars had no field for these items

From Collaborative Stage, they became an entity in Site Specific Factors. These data fields could be used for more than stage and could be extrapolated for studies. Unfortunately, the format was unusable for researchers.

In SSDI, these data were put into multiple fields, in multiple forms – percent positive/range, summary and Allred Score

Step 1

Step 2

Step 3

Step 4

Step 5

Step 6

This information progressed into Collaborative Stage.

The next step was SSDI - Site Specific Disease Items. SSDI format stabilizes the fields that these data are recorded into and provides the data in a format researchers can use.

As these data are collected and used, a determination of their usefulness will be made and further changes or innovations may be made – if warranted.



3







# Commission on Cancer Review of Cancer Registry Data

- Standard 1.6 Cancer Registry Quality Control Plan - Minimum 10% of cases review, physician review
- Standard 1.6 Evaluation of "9" and "99"
- Standard 5.6 Accuracy of Data Call for Data cases must meet 100% compliance
- RQRS Completeness of case
- CP3R Potential to identify trended "holes" in abstraction
- The FUTURE: Standard 6.1 and RCRS!



#### State / Central Registry Edit Sets

- NPCR
- SEER
- State Specific

# Software data base edit sets / validation sets

- Geared to meet the above edits sets
- Required to complete a case

#### **Facility Edit Sets**

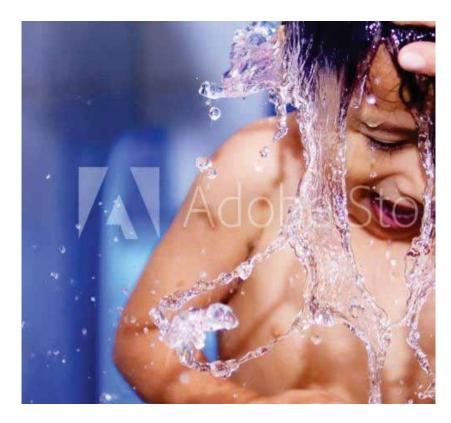


#### National Quality Issues

- Resection of Brain Tumors
- Class of Case physician with admitting privileges
- Grade Coding (before the 2018 changes)

#### Regional Quality Issues

- Network with your peers
- State meeting information a/o central registry issues
- Keep your ears open





#### **New Data Items**

- There is no scarcity
- Do a random review
- Do a sampling review

### Internal Reviews – specific to your facility

- Data specific
- New services
- Individual registrar based



## Solitary Registrar Review

#### Identify your trended issues

- COC Tools
- State/Central Registry
- Completion validation edits
- Run genEdits on a selected dataset

#### Facility Network

Cooperative agreement within network

#### **Outside Sources**

- Cooperative agreement with unaffiliated facility
- Independent Consultant
- Cancer Registry Services

# Multiple Registrar Review

Manager Driven

PEER Driven

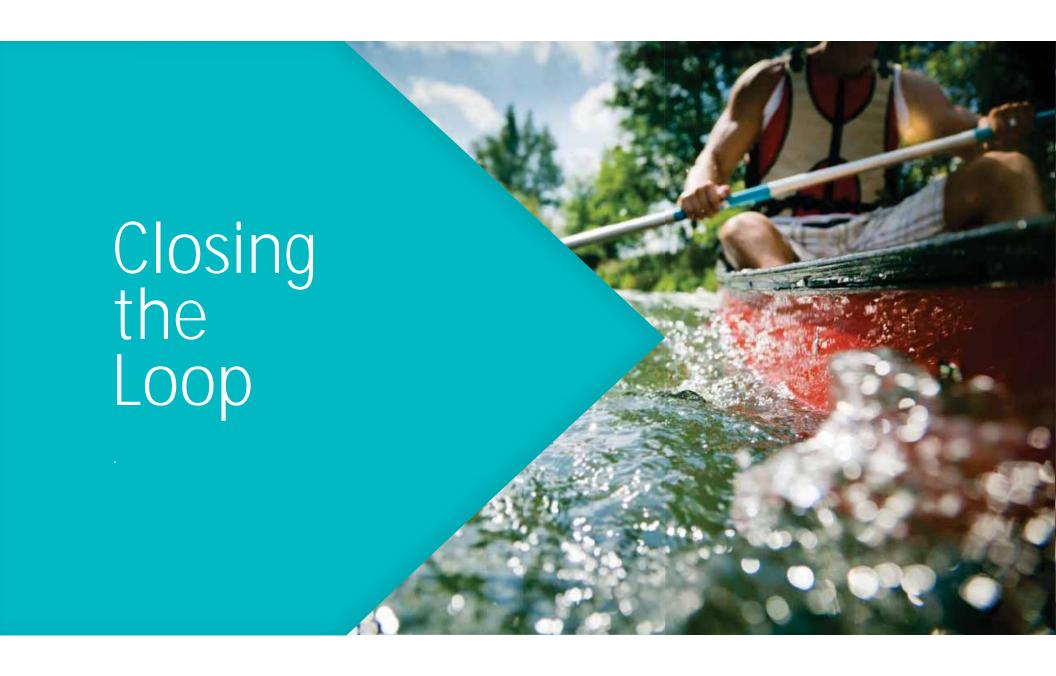
**Facility Network** 

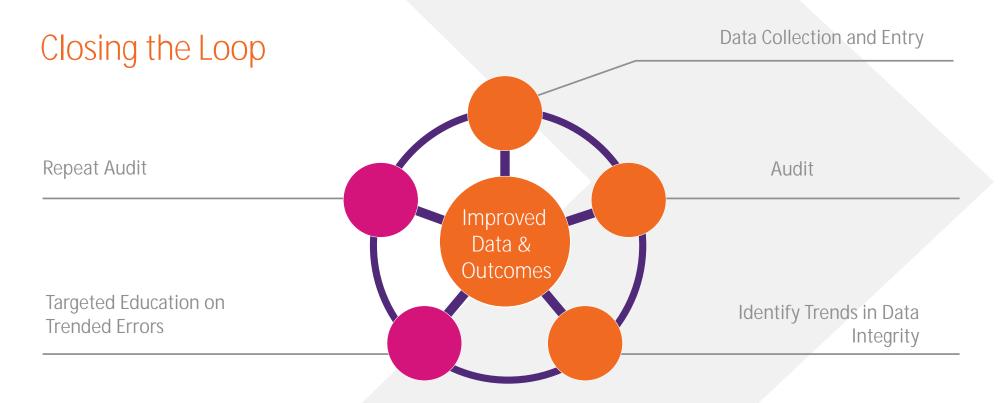
 Cooperative agreement within network

#### **Outside Sources**

- Cooperative agreement with unaffiliated facility
- Independent Consultant
- Cancer Registry Services

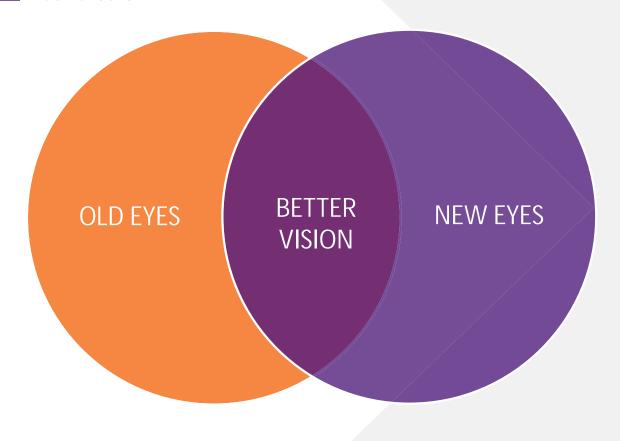








#### CONCLUSION



**OLD EYES** 

+

**NEW EYES** 

=

**BETTER VISION** 

=

**BETTER DATA & OUTCOMES** 





# Questions?

Theresa.Vallerand@cioxhealth.com

910-939-9362



