

## Collaborative Stage Data Collection System (CSv2) Reporting Requirement

### Centers for Disease Control and Prevention, National Program of Cancer Registries (CDC-NPCR)

CDC-NPCR requires the use of Version 2 of the Collaborative Stage Data Collection System (CSv2) for cancer cases diagnosed on or after January 1, 2010. CDC-NPCR requires the collection of CSv2 data items needed to derive SEER Summary Stage, SSFs for Breast, and SSF 25 for applicable sites (schema discriminators). CDC strongly encourages NPCR registries to collect additional CSv2 data items

#### 2010 Required Input Items for all sites:

Item	Item Name
2800	CS Tumor Size
2810	CS Extension
2820	CS Tumor Size/Ext Eval
2830	CS Lymph Nodes
2850	CS Mets at DX

#### Site Specific Factors (for selected primary sites) required to derive SEER Summary Stage :

##### SSF 1 [Lung](#)

Pleura

Retinoblastoma

##### SSF 2 Corpus Adenosarcoma

Corpus Carcinoma

Corpus Sarcoma

##### SSF 3 Prostate

SSF 25 (to direct each site to the correct subgroup discriminator in the algorithm to derive Summary Stage)

#### Site Specific Factors for Breast (not required to derive Summary Stage) but required by CDC-NPCR

**SSF 1** Breast (ERA positive or negative)

**SSF 2** Breast (PRA positive or negative)

**SSF 8** Breast (IHC Value-shows whether or not the cancer cells have HER2 receptors and/or hormone receptors on their surface)

**SSF9** Breast – Her2 IHC Test Interpretation

**SSF10** Breast – Her2 FISH Value

**SSF11** Breast – Her2 FISH Test Interpretation

**SSF12** Breast – Her2 CISH Value

**SSF13** Breast – Her2 CISH Test Interpretation

**SSF14** Breast – Her2Result of Other/Unknown Test