Accrual Data Elements Table with CTRP-Accepted Values for Complete Trials - Include v4.6

Accrual Data Elements for Complete Studies
Descriptions and instructions for Study Subject and Partial Subject demographic data fields

Accrual Data Element	Definition	Required /Conditional		Accepted Values
		Subject	Partial Subject	
Study Subject ID	Enter the unique Patient ID as per the lead organization or the study site where the subject is registered.	R	R	Any numeric or alphanumeric value
Study Subject Birth Date (MM/YYYY)	Enter the subject's month and year of birth in the format `MM/YYYY`.	R		Accrual application format: MM/YYYY REST Services format: YYYY-MM-DD Batch Upload format: YYYYMM
Study Subject Gender	Select the subject's biological sex. If biological sex information is not available, select **Unknown**.	R		Male Female Unspecified Undifferentiated Unknown
Study Subject Race	Select one or more values for race. To select multiple races, select one race, and then press and hold the **CTRL/ CMD** key as you select the other(s).	R		American Indian or Alaska Native Asian Black or African American Native Hawaiian or Other Pacific Islander Not Reported Unknown White
Study Subject Ethnicity	Select a value for ethnicity.	R		Hispanic or Latino Not Hispanic or Latino Not Reported Unknown
Study Subject Country	Select the study subject's country of residence.	R		
Study Subject Zip Code	Enter the study subject's zip code.	C	C	Required if the Study Subject Country is the US, US territories and outlying islands. Must be in a 5 digit or 9 digit (DDDDD-DDDD) format
Registration Date	Enter the date that the subject was registered on the study.	R	R	Accrual application format: MM/DD /YYYY REST Services format: YYYY-MM-DD Batch Upload format: YYYYMMDD
Disease	Click **Look Up**, and follow the instructions in Selecting Diseases for Study Subject Records. Note: For ICD-O-3, CTRP uses a hybrid model with the codes from IACR and the codes in NCI SEER that are not available in IACR.	R		**Partial Subject:** **Disease** is not required. **ICD-O-3:** If **Disease** is reported, **Site** is also required. **Study Subject:** **Disease Code** is required for all trial types. **ICD-O-3: Disease** and **Site** are required.

Site (for ICD-O-3 Disease Codes)	Click **Look Up**, and follow the instructions in Selecting Sites for Study Subject Records Using ICD-O-3 Codes. Site Code information is available at: http://www.iacr.com.fr/index.php?option=com_content&view=category&layout=blog&id=100&Itemid=577 Note: CTRP uses a hybrid model with the codes from IACR and the codes in NCI SEER that are not available in IACR.	C		**Partial Subject:** **Site** is not required. **ICD-O-3:** If **Site** is reported, **Disease** is also required. **Study Subject:** **Site** is required for all trial types. **ICD-O-3: Site** and **Disease** are required.
Participating Site	Select the appropriate site from the drop-down list.	R	R	
Study Subject method of payment	Select the appropriate payment method.			Private Insurance Medicare Medicare and Private Insurance Medicaid Medicaid and Medicare Military or Veterans Sponsored, NOS Military Sponsored (Including CHAMPUS & TRICARE) Veterans Sponsored Self-Pay (No Insurance) No Means of Payment (No Insurance) Managed Care State Supplemental Health Insurance Other Unknown