Copy of Accrual Data Elements Table with CTRP-Accepted Values for Complete Trials - Include v4.8 - ALT 2/15/2024 v2.0

Accrual Data Element	Definition	Required (R) /Conditional (C)		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 Birthdate	Enter the subject's month and year of birth in the format **MM/YYYY**. **Note: The subject's age reported in CTRP cannot be greater than 120 years.**	R		Accrual applicationformat: MM/YYYY RESTServices format: YYYY-MM-DD Batch Upload format: YYYYMM
Study Subject Sex	Select the subject's biological sex. Ifbiological sexinformation 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		CTRP is using the International Standards Organization country codes: https://www.iso.org/obp/ui/#search
Study Subject Zip Code	Enter the study subject's zip code.	C	C	Required if the Study Subject Country is theUS, 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 registeredonthe study.	R	R	User Interface format: MM/DD/YYYY REST 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.
Disease Site	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	