## **Basic Information for Genomics Thailand**

Participant ID	
	([_][_][_][_]])
Text designations that identify gender. Gender is described as the assemblage of properties that distinguish people on the basis of their societal roles. [Explanatory Comment 1: Identification of gender is based upon self-report and may come from a form, questionnaire, interview, etc.]	<ul><li>○ Female</li><li>○ Male</li><li>○ Unknown</li><li>○ Unspecified/Not reported</li></ul>
Birthday (DD-MM-YYYY)	
Date the consent was obtained and participant recruited	
Projects involved in Genomics Thailand	☐ Cancer ☐ Rare disease ☐ Pharmacogenomics ☐ NCD ☐ Infectious Diseases
Is this the patient or the relatives?	<ul><li>Yes</li><li>No</li></ul>
Relationship to the index case	<ul><li>Uncle/Aunt</li><li>Grandparents</li><li>Parents</li><li>Siblings</li></ul>
For cancer and rare disesae, do the patient have a family history?	<ul><li>No</li><li>Yes</li><li>None cancer</li><li>Not rare disease</li><li>Unkonwn</li></ul>
Hospital that recruited the patient	<ul><li>Referred from other hospital</li><li>Hospital in the project</li><li>Unknown</li></ul>
Health policy used by the patient	<ul><li>○ Universal Coverage</li><li>○ Social Security</li><li>○ Government Healthcare</li><li>○ Private/Insurance</li><li>○ No coverage</li></ul>
Biospecimens obtained from the patient	☐ Blood ☐ Surgical tissues ☐ Buccal swab ☐ Skin biopsy

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Has the patient received any genetic test before	<ul> <li>No</li> <li>Yes but positive result not related to the current disesae</li> <li>Yes but got negative result</li> <li>Yes with positive result for the current condition</li> </ul>
For rare disease, do you know the diagnosis of your condition?	<ul><li>○ No</li><li>○ Yes</li><li>○ Not related to the current condition</li></ul>
How many years have you been given the diagnosis (rounding months up to the nearest year)	
Treatment status of the patient	<ul> <li>Never receive treatment</li> <li>Complete treatment</li> <li>Targeted treatment specific to the disease</li> <li>Paliative care</li> <li>Treatment is not applicable</li> </ul>
Choose the organ with disease manifestation	Brain   Eyes   Head and neck   Lungs or respiratory track   Esophagus   Stomach   Small intestine   Large bowel and anus   Liver and bile ducts   Pancreas   Uterus or Prostate   Ovary or Testis   Other internal reproductive organ   Other exteral reproductive organ   Kidney   Urinary track   Breast   Heart muscle   Heart valve   Cardiac conduction system   Arterial disease   Venous system   Lymphatic vessels   Lymph node   Connective tissue   Red Blood Cells   White Blood Cells   White Blood Cells   Platelets   Bone marrow   Skeletal muscles   Bone and joint   Immune system   Peripheral nervous system   Spinal cord   Vertebrae   Ears/Hearing   Skin

# **Germline Testing Indication**

Cancer Type	<ul> <li>○ Breast cancer</li> <li>○ CA Ovary OR Pancreatic Cancer</li> <li>○ Metastatic Prostate Cancer</li> <li>○ Adenomatous Polyps &gt; 10 or Harmatoma &gt; 2</li> <li>○ Colon cancer or endometrial cancer</li> <li>○ Multiple cancers or hereditary cancer syndromes</li> <li>○ Pediatric cancer (&lt; 18 y-o)</li> <li>○ Rare cancer</li> </ul>
Detail of multiple cancers or hereditary cancer syndrome	
Detail of pediatric cancer or rare cancer	
Breast Cancer	
Age with the first breast cancer	
Male breast cancer	○ Yes ○ No
The patient has a triple-negative breast cancer	○ Yes ○ No
Indication for testing in a female patient with breast cancer before the age of 50	<ul> <li>☐ Second primary breast cancer</li> <li>☐ History of breast cancer in first-degree relatives</li> <li>☐ History of pancreatic cancer in first- or second-degree relatives.</li> </ul>
Additional criteria for breast cancer germline testing	<ul> <li>□ At least 2 first-degree relatives with breast cancer, pancreatic cancer, or prostate cancer</li> <li>□ At least 1 first-degree relatives with breast cancer before 50-year-old.</li> <li>□ History of "SARCOMA"</li> <li>□ History of OTHER cancers before 45-year-old</li> <li>□ History of adrenocortical cancer, glioma, or choroid plexus cancer</li> <li>□ At least 1 first/second-degree relatives with cancer before 45-year-old</li> </ul>
Colon Cancer or Endometrial Cancer	
Additional criteria for germline testing for colon cancer or endometrial cancer	<ul> <li>□ Colon cancer or endometrial cancer before 50-year-old</li> <li>□ History of synchronous/metachronous colon/endometrial cancer</li> <li>□ At least 1 first-degree relative with colon/endometrial cancer before 50-year-old</li> <li>□ Abnormal MMR protein or MSI-H in cancer tissue</li> </ul>



# **Minimal Cancer Registry Information**

ICD-10	
International Statistical Classification of Diseases and Related Health Problems 10th Revision (ICD-10)-WHO Version for ;2016 https://icd.who.int/browse10/2016/en#/II	
Vital Status	
The survival state of the person registered on the protocol.	<ul><li>○ Alive</li><li>○ Dead</li><li>○ Unknown</li><li>○ Not Reported</li></ul>
Death Date (DD-MM-YYYY AD)	
Cause of Death	
Text term to identify the cause of death for a patient.	<ul> <li>○ Cancer Related</li> <li>○ Cardiovascular Disorder, NOS</li> <li>○ End-stage Renal Disease</li> <li>○ Infection</li> <li>○ Not Cancer Related</li> <li>○ Renal Disorder, NOS</li> <li>○ Spinal Muscular Atrophy</li> <li>○ Surgical Complications</li> <li>○ Toxicity</li> <li>○ Not Reported</li> <li>○ Unknown</li> </ul>



Extent of the primary cancer based on evidence	
obtained from clinical assessment parameters	Ŏ T1
determined prior to treatment.	Ŏ T1a
	◯ T1a1
	○ T1a2
	O T1b
	○ T1b1
	O T1b2
	O T1c
	○ T1mi
	○ T2 ○ T3-
	○ T2a
	○ T2a1
	○ T2a2
	○ T2b
	○ T2c
	○ T2d
	○ T3
	○ T3a
	○ T3c
	Ŭ T3d
	Ŏ T4
	Ŭ T4a
	O T4b
	O T4c
	O T4d
	○ T4e
	O TX
	○ Ta
	○ Tis
	Tis (DCIS)
	○ Tis (LCIS)
	○ Tis (Paget's)
	○ Unknown
	○ Not Reported/Less Values

Extent of the regional lymph node involvement for the cancer based on evidence obtained from clinical assessment parameters determined prior to treatment.	N0         N0 (i+)         N0 (mol+)         N0 (mol-)         N1         N1a         N1b         N1bI         N1bIII         N1bIV         N1c         N1mi         N2         N2a         N2b         N2c         N3         N3a         N3b         N3c         N4         NX         Unknown         Not Reported
Extent of the distant metastasis for the cancer based on evidence obtained from clinical assessment parameters determined prior to treatment	<ul> <li>M0</li> <li>M1</li> <li>M1a</li> <li>M1b</li> <li>M1c</li> <li>MX</li> <li>cM0 (i+)</li> <li>Unknown</li> <li>Not Reported/Less Values</li> </ul>
The last known state or condition of an individual's neoplasm	<ul> <li>With tumor</li> <li>Distant met recurrence/progression</li> <li>Loco-regional recurrence/progression</li> <li>Biochemical evidence of disease without structura correlate</li> <li>Tumor free</li> <li>Unknown tumor status</li> <li>not reported/Not Allowed To Collect</li> </ul>
Date of the last known status: [last_known_disease_status]	
Evidence supporting progression	<ul><li>X-ray</li><li>Boon scan</li><li>CT/MRI</li><li>PET/PET-CT</li><li>Others</li></ul>

Treatment	
Treatment received by the patients	<ul> <li>No Treatment</li> <li>Unkown</li> <li>Resection</li> <li>Paliative Surgery</li> <li>Single agent chemotherapy</li> <li>Multiple chemotherapy</li> <li>Bone marrow transplant</li> <li>Targeted therapy</li> <li>Immune checkpoint blockade</li> <li>cellular therapy</li> <li>external radiation</li> <li>internal or radionuclide ablation</li> </ul>
The best improvement achieved throughout the entire course of protocol treatment.	<ul> <li>J-Adjuvant Therapy</li> <li>CPD-Clinical Progression</li> <li>CR-Complete Response</li> <li>CRU-Complete Response Unconfirmed</li> <li>DU-Disease Unchanged</li> <li>IMR-Immunoresponse</li> <li>IPD-Immunoprogression</li> <li>MR-Minimal/Marginal Response</li> <li>MX-Mixed Response</li> <li>Non-CR/Non-PD-Non-CR/Non-PD</li> <li>NPB-No Palliative Benefit</li> <li>NR-No Response</li> <li>PA-Palliative Therapy</li> <li>PB-Palliative Benefit</li> <li>PD-Progressive Disease</li> <li>PPD-Pseudoprogression</li> <li>PR-Partial Response</li> <li>PSR-Pseudoresponse</li> <li>RD-Responsive Disease</li> <li>RP-Response</li> <li>SCR-Stringent Complete Response</li> <li>SD-Stable Disease</li> <li>SPD-Surgical Progression</li> <li>TE-Too Early</li> <li>VGPR-Very Good Partial Response</li> </ul>
Regimen used including dose and cycles	
Detail of radiation therapy received by the patients	

# **Additional Cancer Registry Information**

The type of obtained consent from the subject for participation in the study	<ul><li>Consent by Death</li><li>Consent Exemption</li><li>Consent Waiver</li><li>Informed Consent</li></ul>
Number of days between the date used for index and the date the subject consent was obtained for participation in the study.	
The number of days between the date used for index and to the date the patient was lost to follow-up.	
Date of known first cancer usual the pathological diagnosis date	
The patient was unable to be contacted or seen for follow-up information.	<ul><li>Yes</li><li>No</li><li>Unknown</li></ul>
Date Last Follow Up	

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The type of malignant disease (cellular morphology), as categorized by the World Health Organization's (WHO) International Classification of Diseases for Oncology (ICD-O).	<ul> <li>Acinar Cell Neoplasms</li> <li>Adenomas and Adenocarcinomas</li> <li>Adnexal and Skin Appendage Neoplasms</li> <li>Basal Cell Neoplasms</li> <li>Blood Vessel Tumors</li> </ul>
	<ul> <li>Chronic Myeloproliferative Disorders</li> </ul>
	<ul> <li>Complex Épithelial Neoplasms</li> </ul>
	<ul> <li>Complex Mixed and Stromal Neoplasms</li> </ul>
	<ul> <li>Cystic, Mucinous and Serous Neoplasms</li> </ul>
	<ul> <li>Ductal and Lobular Neoplasms</li> </ul>
	<ul><li>Epithelial Neoplasms, NOS</li></ul>
	<ul><li>Fibroepithelial Neoplasms</li></ul>
	Fibromatous Neoplasms
	○ Germ Cell Neoplasms
	○ Giant Cell Tumors
	○ Gliomas
	<ul><li>Granular Cell Tumors and Alveolar Soft Part Sarcomas</li></ul>
	<ul><li>Hodgkin Lymphoma</li></ul>
	Immunoproliferative Diseases
	C Leukemias, NOS
	C Lipomatous Neoplasms
	Lymphatic Vessel Tumors
	Lymphoid Leukemias     Malignant Lymphomas, NOS or Diffuse
	<ul><li>Malignant Lymphomas, NOS or Diffuse</li><li>Mast Cell Tumors</li></ul>
	Mast Cell Tulliors     Mature B-Cell Lymphomas
	Mature T- and NK-Cell Lymphomas
	Meningiomas
	Mesonephromas
	Mesothelial Neoplasms
	Miscellaneous Bone Tumors
	Miscellaneous Tumors
	Mucoepidermoid Neoplasms
	Myelodysplastic Syndromes
	Myeloid Leukemias
	Myomatous Neoplasms
	<ul><li>Myxomatous Neoplasms</li></ul>
	○ Neoplasms, NOS
	<ul> <li>Neoplasms of Histiocytes and Accessory Lymphoid Cells</li> </ul>
	<ul> <li>Nerve Sheath Tumors</li> </ul>
	<ul><li>Neuroepitheliomatous Neoplasms</li></ul>
	Nevi and Melanomas
	Odontogenic Tumors
	Osseous and Chondromatous Neoplasms
	Other Hematologic Disorders
	Other Leukemias
	<ul><li>Paragangliomas and Glomus Tumors</li><li>Plasma Cell Tumors</li></ul>
	Precursor Cell Lymphoblastic Lymphoma
	<ul> <li>Soft Tissue Tumors and Sarcomas, NOS</li> </ul>
	Specialized Gonadal Neoplasms
	Squamous Cell Neoplasms
	Synovial-like Neoplasms
	Thymic Epithelial Neoplasms
	Transitional Cell Papillomas and Carcinomas
	Trophoblastic neoplasms
	O Unknown Not Reported or Not Applicable

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The primary site of disease, as categorized by the World Health Organization's (WHO) International Classification of Diseases for Oncology (ICD-O). This categorization groups cases into general categories. Reference tissue_or_organ_of_origin on the diagnosis node for more specific primary sites of disease.	Accessory sinuses Adrenal gland Anus and anal canal Base of tongue Bladder Bones, joints and articular cartilage of limbs Bones, joints and articular cartilage of other and unspecified sites Brain Breast Bronchus and lung Cervix uteri Colon Connective, subcutaneous and other soft tissues Corpus uteri Esophagus Eye and adnexa Floor of mouth Gallbladder Gum Heart, mediastinum, and pleura Hematopoietic and reticuloendothelial systems Hypopharynx Kidney Larynx Lip Liver and intrahepatic bile ducts Lymph nodes Meninges Nasal cavity and middle ear Nasopharynx Oropharynx Other and ill-defined digestive organs Other and ill-defined sites Other and ill-defined sites Other and ill-defined sites Other and ill-defined sites within respiratory system and intrathoracic organs Other and unspecified female genital organs Other and unspecified female genital organs Other and unspecified parts of biliary tract Other and unspecified parts of biliary tract Other and unspecified parts of fongue Other and unspecified parts of fongue Other and unspecified parts of fongue Other and unspecified parts of tongue Other endocrine glands and related structures Ovary Palate Pancreas Parotid gland Penis
	Peripheral nerves and autonomic nervous system Placenta Prostate gland

Pyriform sinus
Rectosigmoid junction

Retroperitoneum and peritoneum

central nervous system

Spinal cord, cranial nerves, and other parts of

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○ Rectum ○ Renal pelvis

 $\bigcirc \ \mathsf{Stomach}$ Testis ○ Thymus

Ŏ Trachea

Small intestine

Thyroid glandTonsil

Skin

Demographics	
An individual's self-described social and cultural grouping, specifically whether an individual describes themselves as Hispanic or Latino. The provided values are based on the categories defined by the U.S. Office of Management and Business and used by the U.S. Census Bureau.	<ul><li>○ Hispanic or Latino</li><li>○ Not Hispanic or Latino</li><li>○ Unknown</li><li>○ Not reported/Not allowed to collect</li></ul>
An arbitrary classification of a taxonomic group that is a division of a species. It usually arises as a consequence of geographical isolation within a species and is characterized by shared heredity, physical attributes and behavior, and in the case of humans, by common history, nationality, or geographic distribution. The provided values are based on the categories defined by the U.S. Office of Management and Business and used by the U.S. Census Bureau.	<ul> <li>○ Asian</li> <li>○ White</li> <li>○ American Indian or Alaska native</li> <li>○ Black or African American</li> <li>○ Native Hawaiian or Other Pacific Islanders</li> <li>○ Other</li> <li>○ Unknown/not reported</li> <li>○ Not allowed to collect</li> </ul>
Number of days between the date used for index and the date from a person's date of birth represented as a calculated negative number of days.	
A numeric value representing the calendar year in which an individual was born. (Christian Era: 1900 - 2100)	(Christian Era = Buddhist Era - 543)
The yes/no/unknown indicator used to describe whether the patient was premature (less than 37 weeks gestation) at birth.	<ul><li>Yes</li><li>No</li><li>Unknown</li><li>Not Reported</li></ul>
Numeric value used to describe the number of weeks starting from the approximate date of the biological mother's last menstrual period and ending with the birth of the patient.	(Integer only (0-45))

○ Ureter○ Uterus, NOS○ Vagina○ Vulva○ Unknown or Not Reported

# **Additional Follow-up Information**

The source used to determine the patient's cause of death.	<ul> <li>Autopsy</li> <li>Death Certificate</li> <li>Medical Record</li> <li>Social Security Death Index</li> <li>Unknown</li> <li>Not Reported</li> </ul>
Number of days between the date used for index and the date from a person's date of death represented as a calculated number of days.	
Numeric value to represent the year of the death of an individual.	
Last Follow Up	
Time interval from the date of last follow up to the date of initial pathologic diagnosis, represented as a calculated number of days.	
Time interval from the date of last follow up to the date of initial pathologic diagnosis, represented as a calculated number of days.	
Response	
Did the patient has had a new tumor event after initial treatment?	<ul><li>Yes</li><li>No</li><li>unknown</li><li>not reported/Not Allowed To Collect</li></ul>
Date of the first recurrence or progression	
Number of days between the date used for index and the date the patient's disease recurred	
The date that the best response was recorded (DD-MM-YYYY)	
Number of days between the date used for index and the date of the patient was thought to have the best overall response to their disease.	

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# **Additional Diagnosis Information**

Detail of The Tumor	
Date of Initial Pathological Diagnosis	
Text term used to describe the patient's histologic diagnosis, as described by the World Health Organization's (WHO) International Classification of Diseases for Oncology (ICD-O).	
The anatomic site of origin, of the patient's malignant disease, as described by the World Health Organization's (WHO) International Classification of Diseases for Oncology (ICD-O)	
Number of days between the date used for index and the date the patient was diagnosed with the malignant disease.	
Age at the time of diagnosis expressed in number of days since birth. integer, null	
The patient's age (in years) on the reference or anchor date date used during date obfuscation.	
The age of the patient has been modified for compliance reasons. The actual age differs from what is reported. Other date intervals for this patient may also be modified.	<ul><li>○ True</li><li>○ False</li></ul>
The kind of disease present in the tumor specimen as related to a specific timepoint.	<ul> <li>○ primary</li> <li>○ metastasis</li> <li>○ recurrence</li> <li>○ other</li> <li>○ Unknown</li> <li>○ not reported/Not Allowed To Collect</li> </ul>
ICD-O 3	
The third edition of the International Classification of Diseases for Oncology, published in 2000 used principally in tumor and cancer registries for coding the site histology (morphology) of neoplasms. The study of the structure of the cells and their arrangement to constitute tissues and, finally, the association among these to form organs. In pathology, the microscopic process of identifying normal and abnormal morphologic characteristics in tissues, by employing various cytochemical and immunocytochemical stains. A system of numbered categories for the representation of data	

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The third edition of the International Classification of Diseases for Oncology, published in 2000 used principally in tumor and cancer registries for coding the site (topography). The study of the association among these to form organs. A system of numbered categories for the representation of data	
The anatomic site of origin, of the patient's malignant disease, as described by the World Health Organization's (WHO) International Classification of Diseases for Oncology (ICD-O)	
AJCC Tumor Grading	
Numeric value to express the degree of abnormality of cancer cells, a measure of differentiation and aggressiveness	☐ G1 ☐ G2 ☐ G3 ☐ G4 ☐ GX ☐ GB ☐ High Grade ☐ Low Grade ☐ Unknown ☐ Not Reported
Stage group determined from clinical information on the tumor (T), regional node (N) and metastases (M) and by grouping cases with similar prognosis for cancer.	<ul> <li>Stage 0</li> <li>Stage 0is</li> <li>Stage I</li> <li>Stage IA</li> <li>Stage IA1</li> <li>Stage IA2</li> <li>Stage IB</li> <li>Stage IB1</li> <li>Stage IB2</li> <li>Stage II</li> <li>Stage II</li> <li>Stage IIA1</li> <li>Stage IIA1</li> <li>Stage IIIA1</li> <li>Stage IIA2</li> <li>Stage IIB</li> <li>Stage IIIC</li> <li>Stage IIIC</li> <li>Stage IIII</li> <li>Stage IIII</li> <li>Stage IIIIC</li> <li>Stage IIIC1</li> <li>Stage IIIC1</li> <li>Stage IIIC2</li> <li>Stage IIIC2</li> <li>Stage IV</li> <li>Stage Tis</li> <li>Stage X</li> <li>Unknown</li> <li>Not Reported</li> </ul>



Code to represent the defined absence or presence of distant spread or metastases (M) to locations via vascular channels or lymphatics beyond the regional lymph nodes, using criteria established by the American Joint Committee on Cancer (AJCC).	<ul> <li>M0</li> <li>M1</li> <li>M1a</li> <li>M1b</li> <li>M1c</li> <li>M2</li> <li>MX</li> <li>cM0 (i+)</li> <li>Unknown</li> <li>Not Reported</li> </ul>
The codes that represent the stage of cancer based on the nodes present (N stage) according to criteria based on multiple editions of the AJCC's Cancer Staging Manual.	○ N0       (i+)         ○ N0 (mol+)       ○ N0 (mol+)         ○ N0 (mol-)       ○ N1         ○ N1a       ○ N1bl         ○ N1bl       ○ N1bll         ○ N1bll       ○ N1blll         ○ N1bll       ○ N1blll         ○ N1bll       ○ N1bll         ○ N1bll       ○ N1bll         ○ N1mi       ○ N2         ○ N2a       ○ N2b         ○ N2c       ○ N3         ○ N3a       ○ N3b         ○ N3c       ○ N4         ○ N4       ○ NX         ○ Unknown       ○ Not Reported

The extent of a cancer, especially whether the disease has spread from the original site to other parts of the body based on AJCC staging criteria.	Stage 0 Stage 0a Stage 0is Stage I Stage IA Stage IA1 Stage IA2 Stage IB Stage IB1 Stage IB2 Stage IC Stage II Stage IS Stage II Stage IIA Stage IIA1 Stage IIA1 Stage IIA2 Stage IIB Stage IIIC Stage III Stage IIIC Stage IIIIC Stage IIIC Stage IIIC1 Stage IIIC2 Stage IIIC1 Stage IIIC2 Stage IIIC2 Stage IIIC2 Stage IIIC2 Stage IIIC1 Stage IIIC2 Stage IIIC2 Stage IV Stage IVA Stage IVA Stage IVB Stage IVC
	○ Stage IVB

Code of pathological T (primary tumor) to define the size or contiguous extension of the primary tumor (T), using staging criteria from the American Joint Committee on Cancer (AJCC).	<ul> <li>□ T0</li> <li>□ T1</li> <li>□ T1a1</li> <li>□ T1a2</li> <li>□ T1b</li> <li>□ T1b1</li> <li>□ T1b2</li> <li>□ T1c</li> <li>□ T1mi</li> <li>□ T2</li> <li>□ T2a1</li> <li>□ T2a2</li> <li>□ T2b</li> <li>□ T2c</li> <li>□ T2d</li> <li>□ T3</li> <li>□ T3a</li> <li>□ T3b</li> <li>□ T3c</li> <li>□ T3d</li> <li>□ T3d</li> <li>□ T4</li> <li>□ T4a</li> <li>□ T4b</li> <li>□ T4c</li> <li>□ T4d</li> <li>□ T4c</li> <li>□ T4d</li> <li>□ T4e</li> <li>□ TX</li> <li>□ Ta</li> <li>□ Tis</li> <li>□ Tis</li> <li>□ Tis (Paget's)</li> <li>○ Unknown</li> <li>○ Not Reported</li> </ul>
The version or edition of the American Joint Committee on Cancer Staging Handbooks, a publication by the group formed for the purpose of developing a system of staging for cancer that is acceptable to the American medical profession and is compatible with other accepted classifications.	<ul> <li>☐ 1st</li> <li>☐ 2nd</li> <li>☐ 3rd</li> <li>☐ 4th</li> <li>☐ 5th</li> <li>☐ 6th</li> <li>☐ 7th</li> <li>☐ 8th</li> <li>☐ Unknown</li> <li>☐ Not Reported</li> </ul>
Anaplasia	
Was anaplasia present at the time of diagnosis?	<ul><li>Yes</li><li>No</li><li>Unknown</li><li>Not Reported</li></ul>

Text term to signify whether lymphoma B-symptoms are present as noted in the patient's medical record.  The clinical classification of lymphoma, as defined by the Ann Arbor Lymphoma Staging System.  Indicator that identifies whether a patient with mailignant lymphoma has lymphomatous involvement of an extranodal site.  The pathologic classification of lymphoma, as defined by the Ann Arbor Lymphoma Staging System.  Indicator that identifies whether a patient with mailignant lymphoma has lymphomatous involvement of an extranodal site.  Indicator that identifies whether a patient with mailignant lymphoma has lymphomatous involvement of an extranodal site.  Indicator that identifies whether a patient with mailignant lymphoma has lymphomatous involvement of an extranodal site.  Indicator that identifies whether a patient with mailignant lymphoma has lymphomatous involvement of an extranodal site.  Indicator that identifies whether a patient with mailignant lymphoma has lymphomatous involvement of an extranodal site.  Indicator that identifies whether a patient with mailignant lymphoma has lymphomatous involvement of an extranodal site.  Indicator that identifies whether a patient with mailignant lymphoma has lymphomatous involvement of an extranodal site.  Indicator that identifies whether a patient with mailignant lymphoma has lymphomatous involvement of an extranodal site.  Indicator that identifies whether a patient with mailignant lymphoma has lymphomatous involvement of a stage II	The morphologic findings indicating the presence of a malignant cellular infiltrate characterized by the presence of large pleomorphic cells, necrosis, and high mitotic activity in a tissue sample.	<ul> <li>Absent</li> <li>Diffuse</li> <li>Equivocal</li> <li>Focal</li> <li>Present</li> <li>Sclerosis</li> <li>Unknown</li> <li>Not Reported</li> </ul>
present as noted in the patient's medical record.    ONO   Unknown   Not Reported	<b>Ann Arbor Lymphoma Staging System for Lymphor</b>	na
by the Ann Arbor Lymphoma Staging System.  Stage II Stage III Stage IV Unknown Not Reported  Indicator that identifies whether a patient with malignant lymphoma has lymphomatous involvement of an extranodal site.  The pathologic classification of lymphoma, as defined by the Ann Arbor Lymphoma Staging System.  Stage II Stage IV Unknown Not Reported  Burkitt's lymphoma categorization based on clinical features that differ from other forms of the same disease.  Burkitt's lymphoma categorization based on clinical features that differ from other forms of the same disease.  The number that describes the distance, in millimeters, between the upper layer of the epidermis and the deepest point of tumor penetration.		<ul><li>No</li><li>Unknown</li></ul>
malignant lymphoma has lymphomatous involvement of an extranodal site.  The pathologic classification of lymphoma, as defined by the Ann Arbor Lymphoma Staging System.  Stage I Stage II Stage II Stage II Stage IV Unknown Not Reported  Burkitt's lymphoma categorization based on clinical features that differ from other forms of the same disease.  Burkitt's lymphoma categorization based on clinical features that differ from other forms of the same disease.  The number that describes the distance, in millimeters, between the upper layer of the epidermis and the deepest point of tumor penetration.		<ul><li>Stage II</li><li>Stage IV</li><li>Unknown</li></ul>
by the Ann Arbor Lymphoma Staging System.  Stage II Stage IV Unknown Not Reported  Burkitt's lymphoma categorization based on clinical features that differ from other forms of the same disease.  Burkitt's lymphoma categorization based on clinical features that differ from other forms of the same disease.  Endemic Immunodeficiency-associated, adult Immunodeficiency-associated, pediatric Sporadic, adult Sporadic, pediatric Unknown Not Reported  The number that describes the distance, in millimeters, between the upper layer of the epidermis and the deepest point of tumor penetration.	malignant lymphoma has lymphomatous involvement of	○ No ○ Unknown
features that differ from other forms of the same disease.    Immunodeficiency-associated, adult   Immunodeficiency-associated, pediatric   Sporadic, adult   Sporadic, pediatric   Unknown   Not Reported		<ul><li>Stage II</li><li>Stage IV</li><li>Unknown</li></ul>
millimeters, between the upper layer of the epidermis and the deepest point of tumor penetration.	features that differ from other forms of the same	<ul> <li>Immunodeficiency-associated, adult</li> <li>Immunodeficiency-associated, pediatric</li> <li>Sporadic, adult</li> <li>Sporadic, pediatric</li> <li>Unknown</li> </ul>
Child Pugh Classification	millimeters, between the upper layer of the	
	Child Pugh Classification	

The classification used in the prognosis of chronic liver disease, mainly cirrhosis.	<ul> <li>A</li> <li>A5</li> <li>A6</li> <li>B</li> <li>B7</li> <li>B8</li> <li>B9</li> <li>C</li> <li>C10</li> <li>C11</li> <li>C12</li> <li>Unknown</li> <li>Not Reported</li> </ul>
Rectum Cancer	
The number that represents the area of non-peritonealised bare area of rectum, comprising anterior and posterior segments, when submitted as a surgical specimen resulting from excision of cancer of the rectum.	
The staging classification of liver tumors, as defined by the Children's Oncology Group (COG). This staging system specifically describes the extent of the primary tumor prior to treatment.	<ul><li>Stage I</li><li>Stage II</li><li>Stage III</li><li>Stage IV</li><li>Unknown</li><li>Not Reported</li></ul>
The categorization of patients on the basis of prognostic factors per a system developed by Children's Oncology Group (COG). Risk level is used to assign treatment intensity.	<ul><li>○ High Risk</li><li>○ Intermediate Risk</li><li>○ Low Risk</li><li>○ Unknown</li><li>○ Not Reported</li></ul>
The staging classification of renal tumors, as defined by the Children's Oncology Group (COG).	<ul><li>Stage I</li><li>Stage II</li><li>Stage III</li><li>Stage IV</li><li>Unknown</li><li>Not Reported</li></ul>
The classification of rhabdomyosarcoma, as defined by the Children's Oncology Group (COG).	<ul><li> High Risk</li><li> Intermediate Risk</li><li> Low Risk</li><li> Unknown</li><li> Not Reported</li></ul>
The surgical grade of the musculoskeletal sarcoma, using the Enneking staging system approved by the Musculoskeletal Tumor Society (MSTS).	<ul><li> High Grade (G2)</li><li> Low Grade (G1)</li><li> Unknown</li><li> Not Reported</li></ul>
The metastatic stage of the musculoskeletal sarcoma, using the Enneking staging system approved by the Musculoskeletal Tumor Society (MSTS).	

