Delaware Cancer Registry (DCR)

Non-Hospital Reporting of Cancer

Data Collection, Management and Analysis

General Procedures

January 2020

Delaware Cancer Registry
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INTRODUCTION

The Delaware Cancer Registry was established in 1972 to collect and provide accurate and up-to-date information about cancer in the State of Delaware. Since this time, hospitals have reported each case of cancer diagnosed and/or treated at their facility to the registry. **Effective for cases diagnosed on or after January 1, 1996, the law was amended to require reporting by all health care facilities that diagnose or treat cancer patients.** The Delaware Cancer Control Act can be found at the following site: [http://delcode.delaware.gov/title16/c032/index.shtml](http://delcode.delaware.gov/title16/c032/index.shtml) and is included in the appendix of this manual.

The Delaware Cancer Registry is an essential part of our fight against cancer. Data collected is used for the following activities:

- Monitor trends in cancer incidence and mortality by site, geographic area and demographic characteristics of the population
- Guide cancer control program planning and evaluation
- Assist in prioritizing health resource allocations
- Advance clinical, epidemiological, and health services research
- Evaluate cancer cluster reports

In recent years, because an increasing number of cancer patients are receiving diagnostic and treatment services outside of the hospital, the reporting of cancer case information by physician’s offices and other non-hospital facilities has become more crucial for assuring the completeness of information in our state’s central cancer database. Together with the hospitals, laboratories, and ambulatory surgery centers, the physicians of Delaware play a key role in the collection of information. Without your help, we could not answer many of the questions about cancer in Delaware.
WHO IS RESPONSIBLE TO REPORT?

Effective for cases diagnosed on or after January 1, 1996, all health care facilities that diagnose or treat cancer are required to report cancer case information. Chapter 32 of the Cancer Control Act, states:

“Those required to report to the Department occurrences of cancer and benign tumors will include:

(A) Any physician, surgeon, dentist, podiatrist, or other health care practitioners who diagnose or provide treatment for cancer or benign tumors;

(B) The designated representative of any hospital, dispensary, asylum, or other similar public or private institution that diagnose or provide treatment for cancer or benign tumors; and

(C) The designated representative of any laboratory that examines tissue specimens which disclose the existence of cancer or benign tumor”.

The most common types of cancer diagnosed or treated outside a hospital setting include melanoma, noninvasive bladder tumors, small eye tumors, oral or genital tumors, some prostate and breast tumors, tumors in colorectal polyps, lymphoma, leukemia, multiple myeloma, and other bone marrow primaries.

Please contact the Delaware Cancer Registry at (855) 386-6149 if you have any questions
WHAT IS TO BE REPORTED?

1. **Report** neoplasms described with the following terms:
   a. in situ; noninvasive; intraepithelial; noninfiltrating; stage 0
   b. malignant; cancer; malignant neoplasm, carcinoma

2. **Report** benign tumors of the brain and CNS (for diagnoses after January 1, 2004), in any of the following sites:
   - The brain, meninges, spinal cord, cranial nerves, and other parts of the central nervous system, pituitary gland, craniopharyngeal duct, and pineal gland.

3. **Report** cases when the diagnosis is described with terms such as “apparently”, “Appears”, “compatible with”, “comparable with”, “consistent with”, “favors”, “malignant appearing”, “most likely”, “presumed”, “probable”, “suspect (ed)”, “suspicious (for)”. **Do not report** cases described as “possible”, “worrisome”, potentially malignant”, “questionable”, “suggests”, “cannot be ruled out”, “rule out”, “equivocal”. Additional terms for nonmalignant primary intracranial and central nervous system tumors only are “neoplasm” and “tumor” (beginning with 2004 diagnoses and only for C70.0–C72.9, C75.1–75.3).

4. **Report** each primary site cancer separately. Any subsequent diagnosis of or treatment for cancer in another primary site should be reported as a separate case.

5. **Do not report** when a patient has only a history of cancer with no currently active disease.

6. **Do not report**:
   - Basal cell and squamous cell carcinoma of skin, except of genitalia. (effective 1/1/2003)
   - In situ carcinoma of the cervix uteri
   - Cervical intraepithelial neoplasia grade III (CIN III) and
   - Prostatic intraepithelial neoplasia grade III (PIN III).

*Consult Table A in the appendix for additional description of reportable and non-reportable neoplasms and corresponding ICD-0-3 codes. See Table B for specific reportable neoplasms and corresponding ICD-10-CM diagnosis codes. Use SEER Hematopoietic and Lymphoid Neoplasm Database for Hematopoietic and lymphoid reportable neoplasms at https://seer.cancer.gov/seertools/hemelymph/*
When in doubt about whether to report a case, please feel free to contact the Delaware Cancer Registry at (855) 386-6149
HOW TO REPORT?

Please complete the Cancer Reporting Form (CRF) in the appendix for each primary site diagnosed. If more than one cancer is diagnosed simultaneously, please complete a form for each case. Two (2) versions of the CRF are in the appendix. Reporting facilities should submit cancer cases using the standard CRF (Doc. #35-05-02/02/01/19), except surgery centers which are to report using the version subtitled “Ambulatory Surgery Centers (Doc. #35-05-02/02/01/19)”. Send completed form(s) and supporting documentation to:

Delaware Cancer Registry
256 Chapman Road
Oxford Building, Suite 100
Newark, Delaware 19702
Fax: (302) 283-7201
Phone: (855) 386-6149

PATHOLOGY REPORTS OR OTHER SUPPORTING DOCUMENTATION MUST BE ATTACHED TO THE REPORTING FORM

If information is incomplete, a representative from the Delaware Cancer Registry will contact your office to gather the information required to complete case entry into the state system.

TIME PERIOD FOR REPORTING

All cancer cases being reported to the Delaware Cancer Registry must be submitted within 180 days following initial diagnosis and/or first course of treatment. If a case requires longer than 180 days time to yield sufficient information to complete the Cancer Reporting Form, an extension may be granted by phoning the Delaware Cancer Registry at (855) 386-6149.
$100 FINE

As specified in the Delaware Cancer Control Act, any person or entity who violates any provision of this chapter shall be fined $100 for each violation.

### INFORMATION REQUIRED TO COMPLETE CASE REPORT

<table>
<thead>
<tr>
<th>PATIENT IDENTIFICATION</th>
<th>FIRST COURSE OF TREATMENT</th>
<th>FOLLOW-UP</th>
</tr>
</thead>
<tbody>
<tr>
<td>❖ Patient Name</td>
<td>❖ Watchful waiting</td>
<td>❖ Vital status/tumor status</td>
</tr>
<tr>
<td>❖ Social Security Number</td>
<td>❖ Patient refused treatment</td>
<td>❖ Date of last contact or date of death</td>
</tr>
<tr>
<td>❖ Address at Diagnosis</td>
<td>❖ Surgery</td>
<td></td>
</tr>
<tr>
<td>❖ Marital Status</td>
<td>❖ Radiation</td>
<td></td>
</tr>
<tr>
<td>❖ Sex</td>
<td>❖ Chemotherapy</td>
<td></td>
</tr>
<tr>
<td>❖ Race</td>
<td>❖ Other therapy</td>
<td></td>
</tr>
<tr>
<td>❖ Spanish/Hispanic origin</td>
<td></td>
<td></td>
</tr>
<tr>
<td>❖ Date of Birth</td>
<td></td>
<td></td>
</tr>
<tr>
<td>❖ Birthplace</td>
<td></td>
<td></td>
</tr>
<tr>
<td>❖ Usual occupation/industry</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### DIAGNOSIS

| ❖ Date/place of initial diagnosis |
| ❖ PE/scans/scopes/lab            |
| ❖ Operative/pathology findings   |
| ❖ Residual tumor                 |
| ❖ Diagnostic confirmation        |
| ❖ Hospital referred from/to      |

### CANCER INFORMATION

| ❖ Primary site/Histology/Grade (differentiation) |
| ❖ Tumor Size                                   |
| ❖ Extent of Disease/lymph node involvement    |
| ❖ Staging information                         |
APPENDIX
TITLE 16
Health and Safety
PART III
Vital Statistics

CHAPTER 32. CANCER CONTROL ACT

§ 3201. Short title.
This chapter may be cited as the Delaware Cancer Control Act. (62 Del. Laws, c. 334, § 1; 70 Del. Laws, c. 391, § 1.)

§ 3202. Purpose.
The intent of the General Assembly is to require the establishment and maintenance of a cancer registry for the State. This responsibility is delegated to the Department of Health and Social Services, along with the authority to exercise certain powers to implement this requirement. To ensure an accurate and continuing source of data concerning cancer and certain specified tumors of a benign nature, the General Assembly by this chapter requires certain health care practitioners and all hospitals, clinical laboratories and cancer treatment centers within the State to make available to the Department of Health and Social Services information contained in the medical records of patients who have cancer or tumors of a benign nature. It is intended that the product of these efforts will be a central data bank of accurate, precise and current information regarding the subject diseases. (62 Del. Laws, c. 334, § 1; 70 Del. Laws, c. 391, § 1.)

§ 3203. Definitions.
The following words, terms and phrases, when used in this chapter, shall have the meanings ascribed to them in this section, except where the context clearly indicates a different meaning.

(1) "Benign tumor" means any nonmalignant neoplasm, regardless of the tissue of origin, that appears on the American College of Surgeons most recently published annual list of reportable cancers and benign tumors.

(2) "Cancer" means any malignant neoplasm, regardless of the tissue of origin that appears on the American College of Surgeons most recently published annual list of reportable cancers and benign tumors.

(3) "Department" means the State of Delaware Department of Health and Social Services. (62 Del. Laws, c. 334, § 1; 70 Del. Laws, c. 391, § 1.)
§ 3204. Cancer Registry.

The Department shall adopt, promulgate, amend and repeal any rules and regulations that are consistent with law relative to this chapter and necessary to achieve the purpose and requirements of this chapter. These rules and regulations shall include provisions for:

(1) The establishment and maintenance of an up-to-date registry that shall document every occurrence of cancer and of benign tumor in this State;

(2) The establishment of a procedure for reporting to the Department, within 180 days of initial diagnosis or treatment, every occurrence of cancer and of benign tumor in this State. Such procedure shall include the reporting of specified information that the Department deems necessary and appropriate for the recognition, prevention, control or cure of cancer and benign tumors, and shall minimally include the reporting requirements of the National Cancer Data Base established by the American College of Surgeons, along with information regarding the patient's length of residency in Delaware, primary residential address in Delaware and the location and nature of the patient's primary past employment. Those required to report to the Department occurrences of cancer and benign tumors shall include:

   a. Any physician, surgeon, dentist, podiatrist or other health care practitioner who diagnoses or provides treatment for cancer or benign tumors;
   b. The designated representative of any hospital, dispensary, asylum or other similar public or private institution that diagnoses or provides treatment for cancer or benign tumors; and
   c. The designated representative of any laboratory that examines tissue specimens which disclose the existence of cancer or benign tumor;

(3) The establishment of a procedure for the publication and distribution of forms, instructions and notices required by this chapter or necessary to accomplish the purpose of this chapter; and

(4) The establishment of a procedure to obtain follow-up information from those required to report occurrences of cancer and benign tumors pursuant to this chapter. Any follow-up information deemed necessary by the Department shall be submitted to the Department at least 1 time each year by those required to report occurrences of cancer and benign tumors.

This chapter and any rules or regulations issued pursuant to this chapter shall not apply to any person or private institution that, as an exercise of religious freedom, treats the sick or suffering by spiritual means through prayer alone. (62 Del. Laws, c. 334, § 1; 70 Del. Laws, c. 391, § 1; 73 Del. Laws, c. 431, §§ 1, 2.)

§ 3205. Confidentiality of reports.

(a) Any report of an occurrence of cancer or benign tumor made pursuant to this chapter shall not be divulged nor made public in any way that might tend to disclose the identity of the person to whom it relates. However, patient-identifying information may be exchanged among cancer control agencies as authorized by the Department and upon receipt by the Department of satisfactory assurances by those agencies of the preservation of the confidentiality of such information.

(b) No individual or organization providing information to the Department in accordance with this chapter shall be deemed to be, or held liable for, divulging confidential information. (62 Del. Laws, c. 334, § 1; 63 Del. Laws, c. 288, § 1; 70 Del. Laws, c. 149, § 148; 70 Del. Laws, c. 391, § 1.)
§ 3206. Compulsion prohibited.
Nothing in this chapter shall be construed to compel any individual to submit to any medical or public health examination, treatment or supervision. (62 Del. Laws, c. 334, § 1; 70 Del. Laws, c. 391, § 1.)

§ 3207. Violations.
Any person or entity who violates any provision of this chapter shall be fined $100 for each violation. (62 Del. Laws, c. 334, § 1; 70 Del. Laws, c. 391, § 1; 73 Del. Laws, c. 431, § 3.)

§ 3208. Audit and Abstraction of records by department.
(a) Upon request of a person or organization required to report by § 3204 of this title, the Department may audit records and abstract information that is required to be reported.
(b) Any person or organization failing to report as required by this chapter shall permit the Department to audit records and abstract information that is required to be reported.
(c) The Department may charge a fee to be established by regulation to persons and organizations subjected to an audit pursuant to subsection (a) or (b) of this section. Said person or organization shall reimburse the Department. (73 Del. Laws, c. 431, § 3.)

§ 3209. Reserved.
TABLE A.

REPORTABLE NEOPLASMS – TERMS AND ICD-0-3 CODES

<table>
<thead>
<tr>
<th>Cancer Site/Type Terms</th>
<th>ICD-0 3rd Edition Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>NEOPLASMS THAT ARE REPORTABLE TO THE DELAWARE CANCER REGISTRY</strong></td>
<td></td>
</tr>
<tr>
<td>Malignancy <em>(see exclusions in non-reportable section below)</em></td>
<td>Behavior Code “3”</td>
</tr>
<tr>
<td>Malignant neoplasm</td>
<td></td>
</tr>
<tr>
<td>Cancer</td>
<td></td>
</tr>
<tr>
<td>“Carcinoma In Situ” <em>(see exclusions in non-reportable section below)</em></td>
<td>Behavior Code “2”</td>
</tr>
<tr>
<td>“Stage 0”</td>
<td></td>
</tr>
<tr>
<td>“Noninvasive”</td>
<td></td>
</tr>
<tr>
<td>“Intraepithelial”</td>
<td></td>
</tr>
<tr>
<td>“Noninfiltrating”</td>
<td></td>
</tr>
<tr>
<td><em>Includes:</em> Vaginal Intraepithelial Neoplasia, grade III (VAIN III)</td>
<td>C52.__ ; M-8077/2</td>
</tr>
<tr>
<td>Vulvar Intraepithelial Neoplasia, grade III (VIN III)</td>
<td>C51.__ ; M-8077/2</td>
</tr>
<tr>
<td>Anal Intraepithelial Neoplasia, grade III (AIN III)</td>
<td>C21.1 ; M-8077/2</td>
</tr>
<tr>
<td><em>Non-malignant (benign or borderline) primary brain and central nervous system tumors</em> <em>(diagnosed on or after 1/1/2004), in any of the following sites:</em></td>
<td>Behavior Codes: “0” (Benign) or “1” (Borderline)</td>
</tr>
<tr>
<td>Brain</td>
<td>Site Codes:</td>
</tr>
<tr>
<td>Meninges</td>
<td>C71.0 - 71.9</td>
</tr>
<tr>
<td>Spinal cord, cranial nerves, and other parts of the central nervous system</td>
<td>C70.0 - 70.0</td>
</tr>
<tr>
<td>Pituitary gland</td>
<td>C72.0 - 72.9</td>
</tr>
<tr>
<td>Craniohypophyseal duct</td>
<td>C75.1</td>
</tr>
<tr>
<td>Pineal gland</td>
<td>C75.2</td>
</tr>
<tr>
<td>C61.9; M-8148/2</td>
<td>C75.3</td>
</tr>
<tr>
<td><strong>NON-REPORTABLE NEOPLASMS</strong></td>
<td></td>
</tr>
<tr>
<td>The following skin cancers are NOT reportable**:</td>
<td>Site code C44.__ with histology codes 8000-8110</td>
</tr>
<tr>
<td>Basal cell carcinomas of the skin</td>
<td></td>
</tr>
<tr>
<td>Epithelial carcinomas of the skin</td>
<td></td>
</tr>
<tr>
<td>Papillary carcinomas of the skin</td>
<td></td>
</tr>
<tr>
<td>Squamous cell carcinomas of the skin</td>
<td></td>
</tr>
<tr>
<td>The following in situ neoplasms are NOT reportable:</td>
<td>Site Code; Morphology Code</td>
</tr>
<tr>
<td>Carcinoma in situ of the cervix (CIS)</td>
<td>C53.__ ; M-8077/2</td>
</tr>
<tr>
<td>Cervical Intraepithelial Neoplasia grade III (CIN III)</td>
<td>C53.__ ; M-8077/2</td>
</tr>
<tr>
<td>Cervical Intraepithelial Neoplasia with severe dysplasia (CIN III)</td>
<td>C53.__ ; M-8077/2</td>
</tr>
<tr>
<td>Prostatic Intraepithelial Neoplasia grade III (PIN III)</td>
<td>C61.9; M-8148/2</td>
</tr>
</tbody>
</table>

*Including juvenile astrocytoma, pilocytic astrocytoma and piloid astrocytoma

**Note: skin cancers in the genital sites (vagina, clitoris, labium, vulva, prepuce, penis and scrotum) are reportable. Early or evolving melanoma, in situ or invasive is reportable to Delaware.

1 References: International Classification of Diseases for Oncology, 3\textsuperscript{rd} Edition; NAACCR Standards for Cancer Registries, Vol. II; Standards for Oncology Registry Entry STORE 2018

September 19, 2019
**TABLE B**

**ICD-10-CM CASE FINDING CODES FOR REPORTABLE TUMORS**
*(Effective 10/01/2018)*

Cases reportable to the Delaware Cancer Registry include all invasive and in situ malignant neoplasms and specified benign and borderline neoplasms of the brain and CNS.

The following 2018-2019 Comprehensive ICD-10-CM Case Finding Code list is intended to assist reporting facilities in casefinding of reportable neoplasms.

<table>
<thead>
<tr>
<th>ICD-10-CM</th>
<th>Explanation of Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>C00. - C43., C4A.-, C45. - C48., C49.- - C96.-</td>
<td>Malignant neoplasms (excluding category C44 and C49.A), stated or presumed to be primary (of specified site) and certain specified histologies</td>
</tr>
<tr>
<td><strong>NEW for FY2018:</strong> C96.20 Malignant mast cell neoplasm, unspecified C96.21 Aggressive systemic mastocytosis C96.22 Mast cell sarcoma C96.29 Other malignant cell neoplasm</td>
<td></td>
</tr>
<tr>
<td>C44.00, C44.09</td>
<td>Unspecified/other malignant neoplasm of skin of lip</td>
</tr>
<tr>
<td>C44.10 - C44.19-</td>
<td>Unspecified/other malignant neoplasm of skin of eyelid</td>
</tr>
<tr>
<td>C44.13-</td>
<td>Sebaceous cell carcinoma of skin of eyelid, including canthus Note: Effective 10/1/2018</td>
</tr>
<tr>
<td>C44.20-, C44.29-</td>
<td>Unspecified/other malignant neoplasm skin of ear and external auricular canal</td>
</tr>
<tr>
<td>C44.30-, C44.39-</td>
<td>Unspecified/other malignant neoplasm of skin of other/unspecified parts of face</td>
</tr>
<tr>
<td>C44.40, C44.49</td>
<td>Unspecified/other malignant neoplasm of skin of scalp &amp; neck</td>
</tr>
<tr>
<td>C44.50-, C44.59-</td>
<td>Unspecified/other malignant neoplasm of skin of trunk</td>
</tr>
<tr>
<td>C44.60-, C44.69-</td>
<td>Unspecified/other malignant neoplasm of skin of upper limb, incl. shoulder</td>
</tr>
<tr>
<td>C44.70-, C44.79-</td>
<td>Unspecified/other malignant neoplasm of skin of lower limb, including hip</td>
</tr>
<tr>
<td>C44.80, C44.89</td>
<td>Unspecified/other malignant neoplasm of skin of overlapping sites of skin</td>
</tr>
<tr>
<td>C44.90, C44.99</td>
<td>Unspecified/other malignant neoplasm of skin of unspecified sites of skin</td>
</tr>
<tr>
<td>C49.A-</td>
<td>Gastrointestinal Stromal Tumors Note: GIST is only reportable when it is malignant (/3). GIST, NOS (not stated whether malignant or benign) is a /1 and is not reportable.</td>
</tr>
<tr>
<td>D00.- - D09.-</td>
<td>In-situ neoplasms Note: Carcinoma in situ of the cervix (CIN III-8077/2) and Prostatic Intraepithelial Carcinoma (PIN III-8148/2) are not reportable</td>
</tr>
<tr>
<td>D18.02</td>
<td>Hemangioma of intracranial structures and any site</td>
</tr>
<tr>
<td>D32.-</td>
<td>Benign neoplasm of meninges (cerebral, spinal and unspecified)</td>
</tr>
<tr>
<td>D33.-</td>
<td>Benign neoplasm of brain and other parts of central nervous system</td>
</tr>
<tr>
<td>D35.2 - D35.4</td>
<td>Benign neoplasm of pituitary gland, cranioopharyngeal duct and pineal gland</td>
</tr>
<tr>
<td>D42.-, D43.-</td>
<td>Neoplasm of uncertain or unknown behavior of meninges, brain, CNS</td>
</tr>
<tr>
<td>D44.3 - D44.5</td>
<td>Neoplasm of uncertain or unknown behavior of pituitary gland, cranioopharyngeal duct and pineal gland</td>
</tr>
<tr>
<td>D45</td>
<td>Polycythemia vera (9950/3) ICD-10-CM Coding instruction note: Excludes familial polycythemia (C75.0), secondary polycythemia (D75.1)</td>
</tr>
<tr>
<td>D46.-</td>
<td>Myelodysplastic syndromes (9980, 9982, 9983, 9985, 9986, 9989, 9991, 9992)</td>
</tr>
<tr>
<td>ICD-10-CM</td>
<td>Explanation of Code</td>
</tr>
<tr>
<td>-----------</td>
<td>---------------------</td>
</tr>
<tr>
<td>D47.02</td>
<td>Systemic mastocytosis</td>
</tr>
<tr>
<td>D47.1</td>
<td>Chronic myeloproliferative disease (9963/3, 9975/3) ICD-10-CM Coding instruction note: Excludes the following: Atypical chronic myeloid leukemia BCR/ABL-negative (C92.2_) Chronic myeloid leukemia BCR/ABL-positive (C92.1_) Myelofibrosis &amp; Secondary myelofibrosis (D75.81) Myelophthisic anemia &amp; Myelophthisis (D61.82)</td>
</tr>
<tr>
<td>D47.3</td>
<td>Essential (hemorrhagic) thrombocythemia (9962/3) Includes: Essential thrombocytosis, idiopathic hemorrhagic thrombocythemia</td>
</tr>
<tr>
<td>D47.4</td>
<td>Osteomyelofibrosis (9961/3) Includes: Chronic idiopathic myelofibrosis Myelofibrosis (idiopathic) (with myeloid metaplasia) Myelosclerosis (megakaryocytic) with myeloid metaplasia) Secondary myelofibrosis in myeloproliferative disease</td>
</tr>
<tr>
<td>D47.9</td>
<td>Neoplasm of uncertain behavior of lymphoid, hematopoietic and related tissue, unspecified (9970/1, 9931/3)</td>
</tr>
<tr>
<td>D47.2</td>
<td>Neoplasm of uncertain behavior of lymphoid, hematopoietic and related tissue, unspecified (9960/3, 9970/1, 9971/3, 9931/3)</td>
</tr>
<tr>
<td>D49.6, D49.7</td>
<td>Neoplasm of unspecified behavior of brain, endocrine glands and other CNS</td>
</tr>
<tr>
<td>R85.614</td>
<td>Cytologic evidence of malignancy on smear of anus</td>
</tr>
<tr>
<td>R87.614</td>
<td>Cytologic evidence of malignancy on smear of cervix</td>
</tr>
<tr>
<td>R87.624</td>
<td>Cytologic evidence of malignancy on smear of vagina</td>
</tr>
</tbody>
</table>

**Note:** Pilocytic/juvenile astrocytoma M-9421 moved from behavior /3 (malignant) to /1 (borderline malignancy) in ICD-O-3. However, SEER registries will CONTINUE to report these cases and code behavior as /3 (malignant).

**PLEASE CONTACT THE DELAWARE CANCER REGISTRY AT (855) 386-6149 IF YOU HAVE QUESTIONS.**

**Notes**

- **Reportable** diagnoses include juvenile astrocytoma, pilocytic astrocytoma and piloid astrocytoma; behavior is coded as /3 (malignant).
- **Reportable skin cancers include:**
  - Cancers occurring in the skin of genital sites (any histology) -- including vagina, clitoris, vulva, prepuce, penis, and scrotum.
  - Adnexal carcinomas, adenocarcinomas, lymphomas, melanomas, sarcomas and Merkel cell tumors are reportable
- **Non-reportable skin cancers** (primary site C44.--; histology codes 8000-8110) include basal cell carcinoma and squamous cell carcinoma occurring in non-genital sites.
- In situ carcinoma of the cervix uteri is **not reportable**.
- Prostatic intraepithelial neoplasia (PIN III) is **not reportable**.

^ *This* code list incorporates the latest revisions and additions to the International Classification of Diseases (ICD-10-CM). It is available on the SEER website at the following address: [https://seer.cancer.gov/tools/casefinding/fy2019-casefindinglist-icd10cm.pdf](https://seer.cancer.gov/tools/casefinding/fy2019-casefindinglist-icd10cm.pdf)
The Delaware Cancer Registry (DCR) appreciates your cooperation in complying with data submission requirements for reportable diseases. Every attempt is made to streamline the reporting process and to minimize follow-up contacts with reporting facilities. The following instructions are included to clarify those data items that are commonly left blank or incorrectly coded. Please observe the following instructions to avoid additional data request calls from the DCR.

GENERAL INSTRUCTIONS

a) Complete a CRF for each patient your facility diagnosed with or treated for a reportable disease. Do NOT assume that a hospital or other clinician your facility referred a patient to will submit the data.

b) Include pathology/cytology reports with the completed CRF. If these reports are not in your patient records then be sure to note that under COMMENTS.

PRACTITIONER IDENTIFICATION

1. Practitioner/Facility Name – Indicate the name of the attending clinician/facility that is reporting a cancer diagnosis/treatment.

2. Person completing form – Indicate the name of the person completing the CRF.

CASE IDENTIFICATION

1. Sex – Indicate patient’s sex at birth.

2. Race 1, 2 – Indicate the appropriate race group(s) the patient belongs to.

3. Ethnicity – Indicate whether the patient is of Spanish/Latin descent.

4. Patient’s Usu. Occupation - Indicate what job the patient worked for the majority of his/her career, regardless of whether patient is currently retired. For example, if the patient delivered the US mail for 30 years but is now retired then enter “postal carrier” as the occupation, not “retired”.

5. Company or Industry – Indicate the patient’s employer or the kind of business the patient worked in.
CANCER IDENTIFICATION

1. **Date of Initial Diagnosis** – For a specimen sent to pathology, indicate the date the specimen was COLLECTED, not the date that pathology returned a positive diagnosis.

2. **Place of Diagnosis** – Indicate the facility/office where the specimen was collected (e.g., name of physician’s office or ambulatory surgery center).

3. **Primary Cancer Site** – Indicate where the cancer originated (e.g. breast, prostate, bone marrow, skin)

4. **Histology** – Indicate the type of tissue involved (e.g. adenocarcinoma, acute lymphocytic leukemia, melanoma)

5. **Laterality** – Indicate which side of organ is involved with cancer - for example: right breast, skin of mid-back, left and right lung, left kidney.

6. **Diagnostic Confirmation** – Indicate what process/procedure(s) was used to substantiate the cancer diagnosis.

7. **Summary Stage** – Indicate the stage of the cancer.

CANCER DIRECTED 1ST COURSE OF TREATMENT

1. Indicate what treatment(s) the patient has undergone. Be sure to include the date the treatment began.

2. **Type** – Indicate the name of the surgical procedure, drug, or therapy the patient underwent and the amount received. In addition, no treatment is a form of treatment. Be sure to indicate when the treatment plan is either watchful waiting or when the patient refuses treatment.

3. **Date of Last Contact (or Death)** – Indicate when your facility last saw the patient. **If the patient has expired** then provide the date of death and circle the word “Death”.

4. **Evidence of Cancer at Last Visit?** – Indicate whether the patient was cancer free at last visit.

5. **Patient Referred From** – Indicate the name and specialty of the physician that sent the patient to your facility.

6. **Patient Referred To** – Indicate the name and specialty of the physician. Also, provide the name of the facility if applicable.
Instructions:
1. Please type or print clearly.
2. Complete this form for each cancer diagnosed.
3. Mail/fax completed form along with pathology report and any supporting documentation to:
   DELAWARE CANCER REGISTRY
   256 Chapman Road
   Oxford Building, Suite 100
   Newark, DE 19702
   Phone: (855)-386-6149
   Fax: (302) 741-9029

PRACTITIONER IDENTIFICATION
Practitioner/Facility Name: ____________________________________________  Practitioner/facility # ________
Phone: ____________________________  Address: __________________________________________________________
Person completing form: ____________________________  Date Form completed: ____________________________

CASE IDENTIFICATION
Patient’s Last Name: ______________________________________  First Name: ____________________________  MI: _______
Sex: ☐ Male  ☐ Female  Soc. Sec. #: _______ - _______ Date of Birth (MM-DD-YYYY): _______ - _______
Marital Status: ☐ Single  ☐ Married  ☐ Divorced  ☐ Widowed  ☐ Unknown

RACE 1
☐ African American  ☐ Asian (specify)________________________
☐ White  ☐ Other (specify)________________________
☐ American Indian/Alaskan native

RACE 2
☐ African American  ☐ Asian (specify)________________________
☐ White  ☐ Other (specify)________________________
☐ American Indian/Alaskan native

ETHNICITY - Hispanic/Latin Origin: ☐ No ☐ Yes  Specify if yes: ____________________________
Patient’s usual occupation: ____________________________  Company or Industry: ____________________________
Patient’s address at time of diagnosis: Street: ____________________________  City: ____________________________
State: _______  Zip Code: _______  Place of Birth: ____________________________

CANCER DIAGNOSTIC DATA
Date of Initial Diagnosis: ____________________________  Place of Diagnosis (office/facility name): ____________________________
Primary Cancer Site: ____________________________  Histology: ____________________________  Grade: ____________________________
Laterality: ☐ Left  ☐ Mid  ☐ Right  If Melanoma: Ulceration present? ☐ Yes  ☐ No  Tumor Depth: ______ mm

Diagnostic Confirmation (Check all that apply)
☐ Histology/pathology  ☐ Cytology  ☐ Radiology  ☐ Lab Test/Marker Study  ☐ Endoscopy
☐ Immunophenotyping  ☐ Genetic studies  ☐ Clinical diagnosis  ☐ Others (specify) ____________________________
Findings: __________________________________________________________

STAGING
Summary Stage: ☐ In situ  ☐ Localized  ☐ Regional, direct extension  ☐ Regional lymph nodes  ☐ Distant  ☐ Unknown
AJCC Stage: T____ N_____ M_____  Stage _______  Residual Tumor: ____________________________

CANCER DIRECTED FIRST COURSE OF TREATMENT
☐ Watchful Waiting: Date _____________  ☐ Patient Refused TX: Date _____________

<table>
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<tr>
<th>Surgery</th>
<th>Chemotherapy</th>
<th>Radiation Therapy</th>
<th>Hormone Therapy</th>
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Patient status: ☐ Alive  ☐ Dead  ☐ Unknown  Date of last contact (or death): ____________________________
Evidence of cancer at last visit? ☐ Yes  ☐ No  Patient Referred From: ____________________________
Patient Referred To: ____________________________  Comments: ____________________________
DELAWARE CANCER REGISTRY REPORTING FORM for Ambulatory Surgery Centers

Instructions:
1. Print clearly or type.
2. Complete this form for each cancer diagnosed.
3. Mail/fax completed form along with pathology report and any supporting documentation to:

DELAWARE CANCER REGISTRY
256 Chapman Road
Oxford Building, Suite 100
Newark, DE 19702
Phone: (855) 386-6149
Fax: (302) 741-9029

PRACTITIONER IDENTIFICATION

Practitioner Name: ____________________________ Phone: ____________________________
Practitioner Address: __________________________________________________________________
Person completing form: ____________________________ Date Form Completed: ________________

CASE IDENTIFICATION

Patient’s Last Name: ____________________________ First Name: ____________________________ Middle Initial: ________
Sex: □ Male □ Female Social Security #: ______-____-______ Date of Birth: ______-____-______
Marital Status: □ Single □ Married □ Divorced □ Widowed □ Unknown

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<thead>
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<th>Race 2</th>
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<td>□ White</td>
</tr>
<tr>
<td>□ American Indian/ Alaskan native</td>
<td>□ American Indian/ Alaskan native</td>
</tr>
<tr>
<td>□ Other (specify)</td>
<td>□ Other (specify)</td>
</tr>
</tbody>
</table>

ETHNICITY - Hispanic/Latino Origin: □ No □ Yes Specify if yes: ____________________________

Patient’s address at time of diagnosis: Street: __________________________________________
City: ____________________________ State: _______ Zip Code: ____________________________

Patient’s usual occupation: ____________________________ Company or industry: ____________________________

CANCER DIAGNOSTIC DATA (please attach pathology report): Date of Initial Diagnosis: ________________

Place of Diagnosis (office/facility name): ____________________________
Primary Site of Cancer: ____________________________ Histology: ____________________________
Grade: ____________________________ Laterality: □ Left □ Mid □ Right

Patient Referred From: ____________________________
Patient Referred To: ____________________________

If available, please note any additional information on stage of cancer and first course of treatment:

________________________________________________________________________________________
________________________________________________________________________________________
________________________________________________________________________________________
________________________________________________________________________________________
________________________________________________________________________________________

Revised February 2019
Revised February 2020