

# FLORIDA CANCER DATA SYSTEM

## PREFACE

In 1978, the Department of Health and Rehabilitative Services, now known as the Florida Department of Health, contracted with the Sylvester Comprehensive Cancer Center/University of Miami School of Medicine to implement and maintain the Florida Cancer Data System (FCDS). FCDS has been operational and collecting incidence data on cancer cases seen in Florida hospitals on or after January 1, 1981. Ambulatory diagnostic/treatment centers and pathology laboratories began reporting with patients seen on or after July 1, 1997. Dermatologists began reporting with patients seen on or after January 1, 2011. Urologists, Medical Oncologists, and Hematology/Oncologists began reporting patients seen on or after January 1, 2013. Additional specialty physician reporting is expected in the future.

Cancer reporting to FCDS is mandated by Florida statutes. All cancer cases seen in any health facility licensed under Florida Statute Section 395 or Section 408.07 must be reported to FCDS according to Florida Statutes Section 385.202. This includes all hospitals, ambulatory diagnostic and treatment centers, clinical laboratories and physician practices.

Currently, FCDS processes over 185,000 cancer cases each year. When these cases are unduplicated, there are approximately 110,000 newly diagnosed incidence cancer cases per year. Currently, the FCDS database contains approximately 3,500,000 cases.

The 2013 edition of the Florida Physicians' Cancer Reporting Manual is compatible with national reporting standards. These standards are created and endorsed by the Center for Disease Control and Prevention/National Program of Cancer Registries (CDC/NPCR), the North American Association of Central Cancer Registries (NAACCR), the National Cancer Institute/Surveillance Epidemiology & End Results Program (NCI/SEER), and the American College of Surgeons/Commission on Cancer (ACoS/CoC).

## CONFIDENTIALITY

According to Florida Statute 381, Public Health: General Provisions, "Information submitted in reports required by this section is confidential, exempt from the provisions of s.119.07 (1), and is to be made public only when necessary to public health. A report so submitted is not a violation of the confidential relationship between practitioner and patient."

The Health Insurance Portability and Accountability Act of 1996 (HIPAA) became law April 14, 2001. While most organizations had two full years until April 14, 2003 to comply, questions regarding how this new law impacts cancer reporting continues to arise. The North American Association of Central Cancer Registries (NAACCR) has provided materials that address these questions. As you will see, HIPAA regulations only impact current state cancer reporting procedures. Specifically,

**HIPAA allows for the reporting of identifiable cancer data to public health entities. Because the Florida Cancer Data System falls under the definition of a public health entity, HIPAA allows your medical practice to report data to the Florida Department of Health via the Florida Cancer Data System (FCDS) in compliance with Florida State Law and Florida Administrative Code Rule 64D.**

**Written informed consent/release of information from a cancer patient reported to public health entities is not required under HIPAA; you must simply document that the reporting has occurred.**

FCDS continues to adhere to all Florida Statutes and Department of Health Administrative Rules and Guidelines, and follow strict security measures to assure patient and institutional confidentiality.

## IMMUNITY FROM LIABILITY

No institution or individual complying with Florida statutes 385.202, 405.01, 381.0031, and Florida State Administrative Code(may not have latest update) Rules 64D-3.004 and 64D3.034 shall be civilly or criminally liable for divulging information or providing materials to the statewide registry as required by the law.

## FLORIDA STATUTE

Title XXIX  
PUBLIC HEALTH

Chapter 381  
General Provisions

### **381.0031 – Epidemiological Research; Report of Diseases of Public Health Significance to Department**

- (1) The department may conduct studies concerning the epidemiology of diseases of public health significance affecting people in Florida.
- (2) Any practitioner licensed in this state to practice medicine, osteopathic medicine, chiropractic medicine, naturopathy, or veterinary medicine; any hospital licensed under part I of chapter 395; or any laboratory licensed under chapter 483 that diagnoses or suspects the existence of a disease of public health significance shall immediately report the fact to the Department of Health.
- (3) An animal control officer operating under s. 828.27, a wildlife officer operating under s. 379.3311, or an animal disease laboratory operating under s. 585.61 shall report knowledge of any animal bite, diagnosis of disease in an animal, or suspicion of a grouping or clustering of animals having similar disease, symptoms, or syndromes that may indicate the presence of a threat to humans.
- (4) The department shall periodically issue a list of infectious or noninfectious diseases determined by it to be a threat to public health and therefore of significance to public health and shall furnish a copy of the list to the practitioners listed in subsection (2). The list shall be based on the diseases recommended to be nationally notifiable by the Council of State and Territorial Epidemiologists and the Centers for Disease Control and Prevention. The department may expand upon the list if a disease emerges for which regular, frequent, and timely information regarding individual cases is considered necessary for the prevention and control of a disease specific to Florida.
- (5) Reports required by this section must be in accordance with methods specified by rule of the department.
- (6) Information submitted in reports required by this section is confidential, exempt from the provisions of s. 119.07(1), and is to be made public only when necessary to public health. A report so submitted is not a violation of the confidential relationship between practitioner and patient.
- (7) The department may obtain and inspect copies of medical records, records of laboratory tests, and other medical-related information for reported cases of diseases of public health significance described in subsection (4). The department shall examine the records of a person who has a disease of public health significance only for purposes of preventing and eliminating outbreaks of disease and making epidemiological investigations of reported cases of diseases of public health significance, notwithstanding any other law to the contrary. Health care practitioners, licensed health care facilities, and laboratories shall allow the department to inspect and obtain copies of such medical records and medical-related information, notwithstanding any other law to the contrary. Release of medical records and medical-related information to the department by a health care practitioner, licensed health care facility, or laboratory, or by an authorized employee or agent thereof, does not constitute a violation of the confidentiality of patient records. A health care practitioner, health care facility, or laboratory, or any employee or agent thereof, may not be held liable in any manner for damages and is not subject to criminal penalties for providing patient records to the department as authorized by this section.
- (8) The department may adopt rules related to reporting diseases of significance to public health, which must specify the information to be included in the report, who is required to report, the method and time period for reporting, requirements for enforcement, and required followup activities by the department which are necessary to protect public health.
- (9) This section does not affect s. 384.25.

*History.*—s. 2, ch. 29834, 1955; ss. 19, 35, ch. 69-106; s. 67, ch. 77-147; s. 4, ch. 89-311; s. 2, ch. 90-347; s. 15, ch. 91-297; s. 2, ch. 95-188; s. 184, ch. 96-406; s. 175, ch. 97-101; s. 4, ch. 98-151; s. 252, ch. 98-166; s. 8, ch. 2000-367; s. 1, ch. 2012-173; s. 15, ch. 2012-184.

*Note.*—Former s. 381.231.

## FLORIDA STATUTE

Title XXIX  
PUBLIC HEALTH

Chapter 385  
Chronic Diseases

### **385.202 – Statewide Cancer Registry**

(1) Each facility licensed under chapter 395 and each freestanding radiation therapy center as defined in s. 408.07 shall report to the Department of Health such information, specified by the department, by rule, which indicates diagnosis, stage of disease, medical history, laboratory data, tissue diagnosis, and radiation, surgical, or other methods of diagnosis or treatment for each cancer diagnosed or treated by the facility or center. Failure to comply with this requirement may be cause for registration or licensure suspension or revocation.

(2) The department shall establish, or cause to have established, by contract with a recognized medical organization in this state and its affiliated institutions, a statewide cancer registry program to ensure that cancer reports required under this section shall be maintained and available for use in the course of any study for the purpose of reducing morbidity or mortality; and no liability of any kind or character for damages or other relief shall arise or be enforced against any hospital by reason of having provided such information or material to the department.

(3) The department or a contractual designee operating the statewide cancer registry program required by this section shall use or publish said material only for the purpose of advancing medical research or medical education in the interest of reducing morbidity or mortality, except that a summary of such studies may be released for general publication. Information which discloses or could lead to the disclosure of the identity of any person whose condition or treatment has been reported and studied shall be confidential and exempt from the provisions of s. 119.07(1), except that:

- (a) Release may be made with the written consent of all persons to whom the information applies;
- (b) The department or a contractual designee may contact individuals for the purpose of epidemiologic investigation and monitoring, provided information that is confidential under this section is not further disclosed; or
- (c) The department may exchange personal data with any other governmental agency or a contractual designee for the purpose of medical or scientific research, provided such governmental agency or contractual designee shall not further disclose information that is confidential under this section.

(4) Funds appropriated for this section shall be used for establishing, administering, compiling, processing, and providing biometric and statistical analyses to the reporting facilities. Funds may also be used to ensure the quality and accuracy of the information reported and to provide management information to the reporting facilities.

(5) The department may, by rule, classify facilities for purposes of reports made to the cancer registry and specify the content and frequency of the reports. In classifying facilities, the department shall exempt certain facilities from reporting cancer information that was previously reported to the department or retrieved from existing state reports made to the department or the Agency for Health Care Administration. The provisions of this section shall not apply to any facility whose primary function is to provide psychiatric care to its patients.

*History.*—ss. 2, 3, 4, 9, ch. 78-171; s. 5, ch. 82-213; s. 2, ch. 83-234; s. 96, ch. 86-220; s. 1, ch. 90-6; s. 3, ch. 95-188; s. 201, ch. 96-406; s. 190, ch. 97-101; s. 31, ch. 97-237; s. 24, ch. 99-397.

*Note.*—Former s. 381.3812.

## FLORIDA STATUTE

Title XXIX  
PUBLIC HEALTH

Chapter 405  
Medical Information Available For Research

### **405.01 Release of medical information to certain study groups; exemption from liability.**

Any person, hospital, assisted living facility, hospice, sanatorium, nursing or rest home or other organization may provide information, interviews, reports, statements, memoranda, or other data relating to the condition and treatment of any person to research groups, governmental health agencies, medical associations and societies, and in-hospital medical staff committees, to be used in the course of any study for the purpose of reducing morbidity or mortality. No liability of any kind or character for damages or other relief shall arise or be enforced against any person or organization by reason of having provided such information or material, or by reason of having released or published the findings and conclusions of such groups to advance medical research and medical education, or by reason of having released or published generally a summary of such studies.

*History.--s. 1, ch. 65-533; s. 19, ch. 90-344; s. 27, ch. 95-210.*

Title XXIX  
PUBLIC HEALTH

Chapter 405  
Medical Information Available For Research

### **405.02 Limitation on publication of released information.**

Research groups, governmental health agencies, organized medical associations and societies, and in-hospital medical staff committees shall use or publish said material only for the purpose of advancing medical research or medical education in the interest of reducing morbidity or mortality, except that a summary of such studies may be released by any such group for general publication.

*History.--s. 2, ch. 65-533; s. 20, ch. 90-344; s. 244, ch. 96-406.*

Title XXIX  
PUBLIC HEALTH

Chapter 405  
Medical Information Available For Research

### **405.03 Confidentiality.**

In all events, the identity of any person whose condition or treatment has been studied shall be confidential and exempt from the provisions of s. [119.07](#)(1).

*History.--s. 3, ch. 65-533; s. 21, ch. 90-344; s. 245, ch. 96-406.*

## FLORIDA STATUTE

Title XXIX  
PUBLIC HEALTH

Chapter 408  
Health Care Administration

**408.07 Definitions.** As used in this chapter, with exception of ss. 408.031-408.045, the term:

- (1) "Accepted" means that the agency has found that a report or data submitted by a health care facility or a health care provider contains all schedules and data required by the agency and has been prepared in the format specified by the agency, and otherwise conforms to applicable rule or Florida Hospital Uniform Reporting System manual requirements regarding reports in effect at the time such report was submitted, and the data are mathematical reasonable and accurate.
- (2) "Adjusted admission" means the sum of acute and intensive care admissions divided by the ratio of inpatient revenues generated from acute, intensive, ambulatory, and ancillary patient services to gross revenues. If a hospital reports only subacute admissions, then "adjusted admission" means the sum of subacute admissions divided by the ratio of total inpatient revenues to gross revenues.
- (3) "Agency" means the Agency for Health Care Administration.
- (4) "Alcohol or chemical dependency treatment center" means an organization licensed under chapter 397.
- (5) "Ambulatory care center" means an organization which employs or contracts with licensed health care professionals to provide diagnosis or treatment services predominantly on a walk-in basis and the organization holds itself out as providing care on a walk-in basis. Such an organization is not an ambulatory care center if it is wholly owned and operated by five or fewer health care providers.
- (6) "Ambulatory surgical center" means a facility licensed as an ambulatory surgical center under chapter 395.
- (7) "Audited actual data" means information contained within financial statements examined by an independent, Florida-licensed, certified public accountant in accordance with generally accepted auditing standards, but does not include data within a financial statement about which the certified public accountant does not express an opinion or issues a disclaimer.
- (8) "Birth center" means an organization licensed under s. 383.305.
- (9) "Cardiac catheterization laboratory" means a freestanding facility that employs or contracts with licensed health care professionals to provide diagnostic or therapeutic services for cardiac conditions such as cardiac catheterization or balloon angioplasty.
- (10) "Case mix" means a calculated index for each health care facility or health care provider, based on patient data, reflecting the relative costliness of the mix of cases to that facility or provider compared to a state or national mix of cases.
- (11) "Clinical laboratory" means a facility licensed under s. 483.091, excluding: any hospital laboratory defined under s. 483.041(6); any clinical laboratory operated by the state or a political subdivision of the state; any blood or tissue bank where the majority of revenues are received from the sale of blood or tissue and where blood, plasma, or tissue is procured from volunteer donors and donated, processed, stored, or distributed on a nonprofit basis; and any clinical laboratory which is wholly owned and operated by physicians who are licensed pursuant to chapter 458 or chapter 459 and who practice in the same group practice, and at which no clinical laboratory work is performed for patients referred by any health care provider who is not a member of that same group practice.
- (12) "Comprehensive rehabilitative hospital" or "rehabilitative hospital" means a hospital licensed by the agency as a specialty hospital as defined in s. 395.002; provided that the hospital provides a program of comprehensive medical rehabilitative services and is designed, equipped, organized, and operated solely to deliver comprehensive medical rehabilitative services, and further provided that all licensed beds in the hospital are classified as "comprehensive rehabilitative beds" pursuant to s. 395.003(4), and are not classified as "general beds."
- (13) "Consumer" means any person other than a person who administers health activities, is a member of the governing body of a health care facility, provides health services, has a fiduciary interest in a health facility or other health agency or its affiliated entities, or has a material financial interest in the rendering of health services.
- (14) "Continuing care facility" means a facility licensed under chapter 651.
- (15) "Critical access hospital" means a hospital that meets the definition of "critical access hospital" in s. 1861(mm)(1) of the Social Security Act and that is certified by the Secretary of Health and Human Services as a critical access hospital.
- (16) "Cross-subsidization" means that the revenues from one type of hospital service are sufficiently higher than the costs of providing such service as to offset some of the costs of providing another type of service in the hospital. Cross-

subsidization results from the lack of a direct relationship between charges and the costs of providing a particular hospital service or type of service.

(17) "Deductions from gross revenue" or "deductions from revenue" means reductions from gross revenue resulting from inability to collect payment of charges. For hospitals, such reductions include contractual adjustments; uncompensated care; administrative, courtesy, and policy discounts and adjustments; and other such revenue deductions, but also includes the offset of restricted donations and grants for indigent care.

18) "Diagnostic-imaging center" means a freestanding outpatient facility that provides specialized services for the diagnosis of a disease by examination and also provides radiological services. Such a facility is not a diagnostic-imaging center if it is wholly owned and operated by physicians who are licensed pursuant to chapter 458 or chapter 459 and who practice in the same group practice and no diagnostic-imaging work is performed at such facility for patients referred by any health care provider who is not a member of that same group practice.

(19) "FHURS" means the Florida Hospital Uniform Reporting System developed by the agency.

(20) "Freestanding" means that a health facility bills and receives revenue, which is not directly subject to the hospital assessment for the Public Medical Assistance Trust Fund as described in s. 395.701.

(21) "Freestanding radiation therapy center" means a facility where treatment is provided through the use of radiation therapy machines that are registered under s. 404.22 and the provisions of the Florida Administrative Code implementing s. 404.22. Such a facility is not a freestanding radiation therapy center if it is wholly owned and operated by physicians licensed pursuant to chapter 458 or chapter 459 who practice within the specialty of diagnostic or therapeutic radiology.

(22) "GRAA" means gross revenue per adjusted admission.

(23) "Gross revenue" means the sum of daily hospital service charges, ambulatory service charges, ancillary service charges, and other operating revenue. Gross revenues do not include contributions, donations, legacies, or bequests made to a hospital without restriction by the donors.

(24) "Health care facility" means an ambulatory surgical center, a hospice, a nursing home, a hospital, a diagnostic-imaging center, a freestanding or hospital-based therapy center, a clinical laboratory, a home health agency, a cardiac catheterization laboratory, a medical equipment supplier, an alcohol or chemical dependency treatment center, a physical rehabilitation center, a lithotripsy center, an ambulatory care center, a birth center, or a nursing home component licensed under chapter 400 within a continuing care facility licensed under chapter 651.

(25) "Health care provider" means a health care professional licensed under chapter 458, chapter 459, chapter 460, chapter 461, chapter 463, chapter 464, chapter 465, chapter 466, part I, part III, part IV, part V, or part X of chapter 468, chapter 483, chapter 484, chapter 486, chapter 490, or chapter 491.

(26) "Health care purchaser" means an employer in the state, other than a health care facility, health insurer, or health care provider, who provides health care coverage for her or his employees.

(27) "Health insurer" means any insurance company authorized to transact health insurance in the state, any insurance company authorized to transact health insurance or casualty insurance in the state that is offering a minimum premium plan or stop-loss coverage for any person or entity providing health care benefits, any self-insurance plan as defined in s. 624.031, any health maintenance organization authorized to transact business in the state pursuant to part I of chapter 641, any prepaid health clinic authorized to transact business in the state pursuant to part II of chapter 641, any multiple-employer welfare arrangement authorized to transact business in the state pursuant to ss. 624.436-624.45, or any fraternal benefit society providing health benefits to its members as authorized pursuant to chapter 632.

(28) "Home health agency" means an organization licensed under part IV of chapter 400.

(29) "Hospice" means an organization licensed under part VI of chapter 400.

(30) "Hospital" means a health care institution licensed by the Agency for Health Care Administration as a hospital under chapter 395.

(31) "Lithotripsy center" means a freestanding facility that employs or contracts with licensed health care professionals to provide diagnosis or treatment services using electro-hydraulic shock waves.

(32) "Local health council" means the agency defined in s. 408.033.

(33) "Market basket index" means the Florida hospital input price index (FHIPI), which is a statewide market basket index used to measure inflation in hospital input prices weighted for the Florida-specific experience which uses multistate regional and state-specific price measures, when available. The index shall be constructed in the same manner as the index employed by the Secretary of the United States Department of Health and Human Services for determining the inflation in hospital input prices for purposes of Medicare reimbursement.

(34) "Medical equipment supplier" means an organization that provides medical equipment and supplies used by health care providers and health care facilities in the diagnosis or treatment of disease.

(35) "Net revenue" means gross revenue minus deductions from revenue.

(36) "New hospital" means a hospital in its initial year of operation as a licensed hospital and does not include any facility, which has been in existence as a licensed hospital, regardless of changes in ownership, for over 1 calendar year.

(37) "Nursing home" means a facility licensed under s. 400.062 or, for resident level and financial data collection purposes only, any institution licensed under chapter 395 and which has a Medicare or Medicaid certified distinct part used for skilled nursing home care, but does not include a facility licensed under chapter 651.

(38) "Operating expenses" means total expenses excluding income taxes.

(39) "Other operating revenue" means all revenue generated from hospital operations other than revenue directly associated with patient care.

(40) "Physical rehabilitation center" means an organization that employs or contracts with health care professionals licensed under part I or part III of chapter 468 or chapter 486 to provide speech, occupational, or physical therapy services on an outpatient or ambulatory basis.

(41) "Prospective payment arrangement" means a financial agreement negotiated between a hospital and an insurer, health maintenance organization, preferred provider organization, or other third-party payor which contains, at a minimum, the elements provided for in s. 408.50.

(42) "Rate of return" means the financial indicators used to determine or demonstrate reasonableness of the financial requirements of a hospital. Such indicators shall include, but not be limited to: return on assets, return on equity, total margin, and debt service coverage.

(43) "Rural hospital" means an acute care hospital licensed under chapter 395, having 100 or fewer licensed beds and an emergency room, and which is:

(a) The sole provider within a county with a population density of no greater than 100 persons per square mile;

(b) An acute care hospital, in a county with a population density of no greater than 100 persons per square mile, which is at least 30 minutes of travel time, on normally traveled roads under normal traffic conditions, from another acute care hospital within the same county;

(c) A hospital supported by a tax district or subdistrict whose boundaries encompass a population of 100 persons or fewer per square mile;

(d) A hospital with a service area that has a population of 100 persons or fewer per square mile. As used in this paragraph, the term "service area" means the fewest number of zip codes that account for 75 percent of the hospital's discharges for the most recent 5-year period, based on information available from the hospital inpatient discharge database in the State Center for Health Statistics at the Agency for Health Care Administration; or

(e) A hospital designated as a Critical Access Hospital by the Department of Health in accordance with federal regulations and state requirements.

Population densities used in this subsection must be based upon the most recently completed United States census.

(44) "Special study" means a nonrecurring data-gathering and analysis effort designed to aid the agency in meeting its responsibilities pursuant to this chapter.

(45) "Teaching hospital" means any Florida hospital officially affiliated with an accredited Florida medical school which exhibits activity in the area of graduate medical education as reflected by at least seven different graduate medical education programs accredited by the Accreditation Council for Graduate Medical Education or the Council on Postdoctoral Training of the American Osteopathic Association and the presence of 100 or more full-time equivalent resident physicians. The Director of the Agency for Health Care Administration shall be responsible for determining which hospitals meet this definition.

*History.--s. 71, ch. 92-33; s. 75, ch. 92-289; s. 13, ch. 93-129; s. 39, ch. 93-217; s. 17, ch. 95-144; s. 38, ch. 97-103; s. 2, ch. 98-14; s. 2, ch. 98-21; s. 14, ch. 98-89; s. 44, ch. 2000-153; s. 28, ch. 2000-163; s. 2, ch. 2000-227. ch. 2003-258; s. 5, ch. 2005-81; s. 77, ch. 2006-197; s. 10, ch. 2006-261.*

## FLORIDA ADMINISTRATIVE CODE

### Rule 64D-3.029 – Diseases or Conditions to be Reported – See Appendix A.

(1) Diseases or conditions listed in subsection (3) below are of public health significance identified by the Department as of the date of these rules which must be reported by the practitioner, hospital, laboratory, or other individuals via telephone (with subsequent written report within 72 hours, see Rules 64D-3.030-.033, F.A.C.), facsimile, electronic data transfer, or other confidential means of communication to the County Health Department having jurisdiction for the area in which the office of the reporting practitioner, hospital, laboratory or patient’s residence is located consistent with the specific section and time frames in subsection (3) below relevant to the practitioners, hospitals and laboratories, respectively. Reporters are not prohibited from reporting diseases or conditions not listed by rule.

(2) Definitions to be used with subsection (3) below:

(a) “*Notifiable Diseases or Conditions*” – The definitions of “suspected case” and “confirmed case” for reportable diseases or conditions are set forth in “Surveillance Case Definitions for Select Reportable Diseases in Florida,” August 2008, incorporated by reference, available online at: [http://www.doh.state.fl.us/disease\\_ctrl/epi/surv/CaseDefAug2008.pdf](http://www.doh.state.fl.us/disease_ctrl/epi/surv/CaseDefAug2008.pdf).

(b) “*Suspect Immediately*” – A notifiable condition of urgent public health importance. Report without delay upon the occurrence of any of the following: Initial suspicion, receipt of a specimen with an accompanying request for an indicative or confirmatory test, findings indicative thereof, or suspected diagnosis. Reports that cannot timely be made during the County Health Department business day shall be made to the County Health Department after-hours duty official. If unable to do so, the reporter shall contact the Florida Department of Health after-hours duty official at (850) 245-4401.

(c) “*Immediately*” – A notifiable condition of urgent public health importance. Report without delay upon the occurrence of any of the following: An indicative or confirmatory test, findings indicative thereof, or diagnosis. Reports that cannot timely be made during the County Health Department business day shall be made to the County Health Department after-hours duty official. If unable to do so, the reporter shall contact the Florida Department of Health after-hours duty official at (850) 245-4401.

(d) “*Next Business Day*” – Report before the closure of the County Health Department’s next business day following suspicion or diagnosis.

(e) “*Other*” – Report consistent with the instruction in and footnotes to subsection (3) below.

(3) “*Table of Notifiable Diseases or Conditions to Be Reported*”.

<b>TABLE OF NOTIFIABLE DISEASES OR CONDITIONS</b> <b>(Excerpts Pertaining to the Reporting of Cancer)</b> <b>To view the entire TABLE OF NOTIFIABLE DISEASES OR CONDITIONS – See Appendix A</b>
Cancer (except non-melanoma skin cancer, and including benign and borderline intracranial and CNS tumors)
Pathological or tissue diagnosis of cancer (except non-melanoma skin cancer and including benign and borderline intracranial and CNS tumors)



## FLORIDA ADMINISTRATIVE CODE

### Rule 64D-3.034 – Cancer Reporting.

- (1) Reporting Requirements:
  - a. Each facility and laboratory licensed under Chapters 395 and 483, and Section 408.07(20), F.S., respectively and practitioners licensed under Chapter 458, 459, 464, F.S., are required to report to the Florida Cancer Data System as required by Section 385.202, F.S., within six (6) months of each diagnosis and within six (6) months of the date of each treatment.
  - b. Each facility shall submit each cancer case report electronically. Those facilities with fewer than 35 cancers annually requiring abstracting may submit paper copies or portions of the medical record, provided the copies contain all of the required information as per (1)(c).
  - c. The data items, coding schemes, definitions, record layouts, and reporting procedures are to follow the guidance provided in the Florida Cancer Data System Data Acquisition Manual (2005, or current edition), incorporated by reference, available at <http://www.fcds.med.miami.edu/inc/downloads.shtml>.
- (2) Notwithstanding subsection (1), each facility, center, and laboratory that reports cancer cases to the Florida Cancer Data System shall make its records available for on-site review by the department or its authorized representatives.

*Rulemaking Authority 381.0011(2), 381.003(2), 381.0031(8), 384.33, 385.202(5), 392.66 FS. Law Implemented 381.0011, 381.003, 381.0031, 384.25, 385.202, 392.53 FS. History—New 11-20-06.*

## FLORIDA ADMINISTRATIVE CODE

### Rule 64D-3.003 – Notification by Laboratories

(1) Each laboratory director or designee in charge of a laboratory shall report, or cause to be reported evidence suggestive of or diagnostic of diseases or conditions listed in subsection 64D-3.002(1), F.A.C., from any specimen derived from a human body, or from an animal in the case of rabies or plague testing, to the county health department director or administrator or the State Health Officer or to either of their designated representatives. Such reports shall be made within 72 hours of recognition by telephone, or other electronic means, or in writing, except for certain specified diseases as indicated by a (T), which shall be reported immediately by telephone and followed by a written report. Exceptions to laboratory reporting as defined by this rule are provided for sexually transmitted diseases including AIDS, as indicated in Rule 64D-3.017, F.A.C.

(2) All reports of cancer identified by laboratories licensed under Chapter 483, F.S., shall be submitted to the Florida Cancer Data System within six (6) months of diagnosis.

(3) The State Health Officer shall periodically, but no less than annually, issue a listing of laboratory test results that are to be reported. The July 1999 “Reportable Laboratory Findings,” incorporated by reference in this rule, shall be updated to reflect changes in technology and practice and may be obtained from the Department of Health, Bureau of Epidemiology, 4052 Bald Cypress Way, Bin A-12, Tallahassee, Florida 32399-1720.

(4) To allow follow-up of laboratory findings by the local county health department director/administrator or their designee, all specimens submitted for laboratory tests or examinations related to a disease or condition listed in subsection 64D-3.002(1), F.A.C., shall be accompanied by certain identifying information. In addition to the name and date of birth of the person from whom the specimen was obtained; the name, address and telephone number of the processing clinical laboratory; and the diagnostic test(s) performed, specimen type and result, the following information shall be provided:

(a) Address, telephone number, race, sex, and ethnicity of the person from whom the specimen was obtained or, if this is not available,

(b) Name, address and telephone number of the submitting physician, health care provider or other authorized person who submitted the specimen.

(5) The practitioner who first authorizes, orders, requests or submits a specimen shall be responsible for obtaining and providing the information required in (4) above at the time the specimen is sent to or received by the laboratory.

(6) Notification of test results shall be submitted by telephone, or other electronic means, or in writing on a form furnished by the laboratory. Reports shall be made within 72 hours of a test result. Any preliminary telephone communication must be followed up by a written report.

(7) If the laboratory that makes the positive finding received the specimen from another laboratory, the laboratory making the positive finding shall be responsible for reporting such results as defined in subsection 64D-3.003(1), F.A.C.

(8) In addition to the reporting requirements pursuant to subsection 64D-3.003(1), F.A.C., each laboratory that obtains a human isolate of *Escherichia coli* O157:H7, or *Neisseria meningitidis* or *Haemophilus influenzae* from a sterile site or *Staphylococcus aureus* with a vancomycin minimum inhibitory concentration (MIC) = or > 8 micrograms per milliliter from any site shall retain a subculture of the isolate on suitable media for at least six months after receipt of the specimen in the laboratory. In lieu of retaining this subculture, the laboratory is permitted to send the subculture to the Florida Department of Health State Central Laboratory, which will maintain a record indicating the date that these subcultures were submitted to the Central Laboratory.

(9) In addition to the reporting requirements pursuant to subsection 64D-3.003(1), F.A.C., each laboratory that makes a finding, or suggestive finding, of malaria or cyclospora parasites in a specimen of a patient shall retain a stained permanent slide for at least six months after receipt of the specimen in the laboratory. In lieu of retaining the slide(s), the laboratory may send such slide(s) to the State of Florida Department of Health Central Laboratory, which will maintain a record indicating the date that these specimens were submitted to the Central Laboratory.

(10) Each laboratory licensed to perform tests for any reportable disease or condition shall make its records for such diseases or conditions available for on-site inspection by the department or its authorized representatives.

(11) Persons submitting specimens for reportable laboratory tests to the Florida Department of Health, pursuant to subsection 64D-3.003(4), F.A.C., are required to supply the laboratories with sufficient information to comply with the provisions of this section.

*Specific Authority 381.0011(13), 381.003(2), 381.0031(6), 384.33 FS. Law Implemented 381.0011, 381.003, 381.0031, 384.25 FS. History—New 12-29-77, Amended 6-7-82, Formerly 10D-3.66, Amended 2-26-92, 7-21-96, Formerly 10D-3.066, Amended 11-2-98, 7-5-99, 6-4-00, 6-9-03. Repealed 11-20-06...*

*Editorial Note: See 64D-3.031*

## FLORIDA ADMINISTRATIVE CODE

### Rule 64D-3.006 – Reports, Medical Facilities and Freestanding Radiation Therapy Centers.

(1) The chief administrative officer of each civilian facility licensed under Chapter 395, F.S., and freestanding radiation therapy centers, as defined in Section 408.07, F.S., shall (and the United States military and Veterans Administration hospitals are requested to) appoint an individual from the staff, hereinafter referred to as “reporting officer,” who shall be responsible for reporting cases or suspect cases of diseases on the notifiable disease list in persons admitted to, attended to, or residing in the facility (cf. Notification by Laboratories, Rule 64D-3.003, F.A.C.).

(2) Reporting of a case or suspected case of notifiable disease or condition by a facility or center fulfills the requirements of the licensed practitioner to report; however, it is the responsibility of the practitioner to ensure that the report is made as stipulated in Rule 64D-3.002, F.A.C. Reports shall be made within 72 hours of diagnosis. Special provisions for reporting sexually transmissible diseases, including HIV infection, are found in Rule 64D-3.016, F.A.C., and for cancer, in subsection 64D-3.006(3), F.A.C.

(3) Reporting of cancer cases by a licensed practitioner, a hospital facility licensed under Chapter 395, F.S., and freestanding radiation therapy centers, as defined in Section 408.07, F.S., to the Florida Cancer Data System as required by Section 385.202, F.S., shall be accomplished within six (6) months of the date of each diagnosis and within six (6) months of the date of each treatment.

(4) Florida Cancer Data System staff will provide each freestanding ambulatory surgical center with an annual list of cancer cases for which reports are required and allow three (3) months from the date of notification for submission of reports to the Florida Cancer Data System for each case on the list. This annual list will be generated by comparing the ambulatory patient data maintained by the Agency for Health Care Administration with the Florida Data System file for each calendar year. This comparison will be made each year after the Florida Cancer Data System file for each year is complete, including all hospital and pathology laboratory data expected for that year. The list sent to each freestanding ambulatory surgical center will contain only those records from the Agency for Health Care Administration ambulatory patient dataset or from cancer case data received from ambulatory centers that cannot be matched with any previously reported case.

(5) For reportable cancer cases, each facility licensed under chapter 395, F.S., and each freestanding radiation therapy center as defined in Section 408.07, F.S., shall electronically submit to the Florida Cancer Data System all available data items as specified in the Data Acquisition Manual and Confidential Abstract Report. Those facilities and centers with fewer than thirty-five (35) cancer cases annually requiring abstracting may submit to FCDS paper copies of portions of the case record that include all available information that is needed for abstracting by FCDS staff. The coding schemes, record layouts, and definitions for these items are those issued by the Florida Cancer Data System in its Data Acquisition Manual and Confidential Abstract Report, DOH Form 2029, dated July 1997, incorporated herein by reference. These documents are available from the Florida Department of Health, Bureau of Epidemiology, 4052 Bald Cypress Way, Bin A-12, Tallahassee, Florida 32399-1720.

*Specific Authority 381.0011(13), 381.003(2), 381.0031(6), 384.33, 385.202(5), 392.66 FS. Law Implemented 381.0011, 381.003, 381.0031, 384.25, 385.202, 392.53 FS. History—New 12-29-77, Amended 6-7-82, Formerly 10D-3.77, Amended 2-26-92, 7-21-96, Formerly 10D-3.077, Amended 11-2-98, 7-5-99, 6-4-00.*

## FLORIDA ADMINISTRATIVE CODE

### Rule 64D-3.030 – Notification by Practitioners.

(1) Each practitioner licensed under Chapters 458, 459, 460, 462, 464, 467 and 474, F.S., and medical examiner appointed pursuant to Chapter 406, F.S., who diagnoses, treats or suspects a case, or who suspects an occurrence of a disease or condition listed in the Table of Notifiable Diseases or Conditions to Be Reported, Rule 64D-3.029, F.A.C., including in persons who at the time of death were so affected, shall report or cause to be reported all such diagnoses or suspicions per this rule. Reporting of specimen results by a laboratory to a county health department director, administrator or designee does not nullify the practitioner's obligation to report said disease or condition.

(2) Any request for laboratory test identification shall be considered a suspicion of disease. However, practitioners need only to report suspected cases if indicated in the "suspect immediately" column under practitioners in the Table of Notifiable Diseases or Conditions to Be Reported, Rule 64D-3.029, F.A.C.

(3) Any report of a notifiable disease or condition required by this rule, except for cancer, congenital anomalies and HIV/AIDS, shall be reported on the Florida Department of Health Disease Report Form (DH Form 2136, 3/06), incorporated by reference, available at the Department of Health, Division of Disease Control, 4052 Bald Cypress Way, Bin A-09, Tallahassee, FL 32399-1714, or on a form supplied by the provider that includes the following:

(a) The patient's:

1. First and last name, including middle initial;
2. Address, including city, state and zip code;
3. Telephone number, including area code;
4. Date of birth;
5. Sex;
6. Race;
7. Ethnicity (specify if of Hispanic descent or not of Hispanic descent);
8. Pregnancy status if applicable;
9. Social Security number;
10. Date of onset of symptoms;
11. Diagnosis.

(b) Type of diagnostic tests (for example culture, IgM, serology, Mantoux TB skin test, nucleic acid amplification test or Western Blot);

(c) Type of specimen (for example stool, urine, blood, mucus, etc.);

(d) Date of specimen collection;

(e) Site (for example cervix, eye, etc., if applicable);

(f) Diagnostic test results including: reference range, titer when quantitative procedures are performed, and all available results concerning additional characterization of the organism;

(g) For Tuberculosis, the 15-digit spoligotype (octal code) must be reported;

(h) Treatment given;

(i) Name, address and telephone number of the attending practitioner;

(j) Other necessary epidemiological information as well as additional specimen collection or laboratory testing requested by the county health department director or administrator or their designee.

(4) The practitioner who first authorizes, orders, requests or submits a specimen to a licensed laboratory for testing for any agent listed in Rule 64D-3.029, F.A.C., shall obtain and provide the information required by subparagraphs 64D-3.031(3)(a)1.-9., F.A.C., at the time the specimen is sent.

(5) Special reporting requirements for HIV and AIDS:

(a) All cases of HIV or AIDS, which meet the Centers for Disease Control and Prevention (CDC) case definitions set forth in CDC Guidelines for National Human Immunodeficiency Virus Case Surveillance, Including Monitoring for Human Immunodeficiency Virus Infection and Acquired Immunodeficiency Syndrome, published in Morbidity and Mortality Weekly Report (MMWR) Vol. 48 [RR-13, December 10, 1999], incorporated by reference, available online at: [www.cdc.gov/mmwr/PDF/RR/RR4813.pdf](http://www.cdc.gov/mmwr/PDF/RR/RR4813.pdf), shall be reported on the Adult HIV/AIDS Confidential Case Report, CDC 50.42A Rev. 03/2007, incorporated by reference, or the Pediatric HIV/AIDS Confidential Case Report, CDC 50.42B Rev. 01/2003, incorporated by reference, along with the Department of Health Addendum for Adult HIV/AIDS

Confidential Case Report, DH Form 2134, (09/08), incorporated by reference. All forms are available at county health departments or at the Department of Health, Bureau of HIV/AIDS, 4052 Bald Cypress Way, Bin A-09, Tallahassee, Florida 32399-1715, (850) 245-4334.

(b) HIV exposed newborns shall be reported on the Pediatric HIV/AIDS Confidential Case Report, CDC 50.42B Rev. 01/2003, incorporated by reference in paragraph 64D-3.030(5)(a), F.A.C.

(6) Each practitioner who makes a diagnosis of or treats any notifiable disease or condition shall make their patient medical records for such diseases or conditions available for on-site inspection by the Department or its authorized representatives.

*Rulemaking Authority 381.0011(2), 381.003(2), 381.0031(7), (8), 383.06, 384.25(1), 384.33, 392.53(1), 392.66 FS. Law Implemented 381.0011(3), 381.003(1), 381.0031(2), (4), (8), 384.23, 384.25, 385.202, 392.53 FS. History—New 11-20-06, Amended 11-24-08.*

## PUBLIC LAW 102-515

Public Law 102-515 – Oct. 24, 1992  
Public Law 102-515 102d Congress  
**106 STAT. 3372**

**An Act** Entitled the “Cancer Registries Amendment Act”.

*Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,*

### **SECTION 1. SHORT TITLE.**

This Act may be cited as the “Cancer Registries Amendment Act”.

### **SEC. 2. FINDINGS AND PURPOSE.**

(a) **FINDINGS.**—Congress finds that— (1) cancer control efforts, including prevention and early detection, are best addressed locally by State health departments that can identify unique needs; (2) cancer control programs and existing statewide population-based cancer registries have identified cancer incidence and cancer mortality rates that indicate the burden of cancer for Americans is substantial and varies widely by geographic location and by ethnicity; (3) statewide cancer incidence and cancer mortality data, can be used to identify cancer trends, patterns, and variation for directing cancer control intervention; (4) the American Association of Central Cancer Registries (AACCR) cites that of the 50 States, approximately 38 have established cancer registries, many are not statewide and 10 have no cancer registry; and (5) AACCR also cites that of the 50 States, 39 collect data on less than 100 percent of their population, and less than half have adequate resources for insuring minimum standards for quality and for completeness of case information. (b) **PURPOSE.**—It is the purpose of this Act to establish a national program of cancer registries.

### **SEC. 3. NATIONAL PROGRAM OF CANCER REGISTRIES.**

Title III of the Public Health Service Act (42 U.S.C. 241 et seq.) is amended by adding at the end the following new part: “PART M—NATIONAL PROGRAM OF CANCER REGISTRIES  
42 USC 280e. “**SEC. 399H. NATIONAL PROGRAM OF CANCER REGISTRIES.**

“(a) **IN GENERAL.**—The Secretary, acting through the Director of the Centers for Disease Control, may make grants to States, or may make grants or enter into contracts with academic or nonprofit organizations designated by the State to operate the State’s cancer registry in lieu of making a grant directly to the State, to support the operation of population-based, statewide cancer registries in order to collect, for each form of in-situ and invasive cancer (with the exception of basal cell and squamous cell carcinoma of the skin), data concerning—

## PUBLIC LAW 107-260

Public Law 107-260 – Oct. 29, 2002  
Public Law 107-260 107<sup>th</sup> Congress  
116 STAT. 1743

An Act to amend the Public Health Service Act to provide for the collection of data on benign brain-related tumor through the national program of cancer registries.

*Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,*

### SECTION 1. SHORT TITLE.

This Act may be cited as the “Benign Brain Tumor Cancer Registries Amendment Act”.

### SEC. 2. NATIONAL PROGRAM OF CANCER REGISTRIES; BENIGN BRAINRELATED TUMORS AS ADDITIONAL CATEGORY OF DATA COLLECTED.

- (a) In GENERAL—Section 399B of the Public Health Service Act (42 U.S.C. 280e), as redesignated by section 502 (2) (A) of Public Law

106-310 (114 Stat. 1115), is amended in subsection (a)—

- (1) by redesignating paragraphs (1) through (5) as subparagraphs (A) through (3), respectively, and indenting appropriately;
- (2) by striking “(a) IN GENERAL—The Secretary” and inserting the following:

- (a) IN GENERAL—

“(1) STATEWIDE CANCER REGISTRIES—The Secretary”;

- (3) in the matter preceding subparagraph (A) (as so redesignated). By striking “population-based” and all that follows through “data” and inserting the following: “population-based, statewide registries to collect, for each condition specified in paragraph (2)(A), data”; and

- (4) by adding at the end the following:

“(2) CANCER; BENIGN BRAIN-RELATED TUMORS—

“(A) IN GENERAL—For purposes of paragraph (1), the conditions referred to in this paragraph are the following:

“(i) Each form of in-situ and invasive cancer with the exception of basal cell and squamous cell carcinoma of the skin), including malignant brain-related tumors.

“(ii) Benign brain-related tumors

“(B) BRAIN-RELATED TUMOR—For purposes of subparagraph (A):

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“(i) The term ‘brain-related tumor’ means a listed primary tumor (whether malignant or benign) occurring in any of the following sites:’

“(I) The brain, meninges, spinal cord, cauda equina, a cranial nerve or nerves or any other part of the central nervous system.

“(II) The pituitary gland, pineal gland, or craniopharyngeal duct.

“(ii) The term ‘listed’, with respect to a primary tumor, means a primary tumor that is listed in the International Classification of Diseases for Oncology (commonly referred to as the ICD-O).

“(iii) The term ‘International Classification of Diseases for Oncology’ means a classification system that includes topography (site) information and histology (cell type information) developed by the World Health Organization, in collaboration with international centers, to promote international comparability in the collection, classification, processing and presentation of cancer statistics. The ICDO system is a supplement to the International Statistical Classification of Diseases and Related Health Problems (commonly known as the ICD) and is the standard coding system used by cancer registries worldwide. Such term includes any modification made to such system for purposes of the United States. Such term further includes any published classification system that is internationally recognized as a successor to the classification system referred to in the first sentence of this clause.

“(C) STATEWIDE CANCER REGISTRY—References in this section to cancer registries shall be considered to be references to registries described in this subsection.”

- (b) APPLICABILITY—The amendments made by subsection (a) apply to grants under section 399B of the Public Health Service Act for fiscal year 2002 and subsequent fiscal years, except that, in the case of a State that received such a grant for fiscal year 2000, the Secretary of Health and Human Services may delay the applicability of such amendments to the State for not more than 12 months if the Secretary determines that compliance with such amendments requires the enactment of a statute by the State or the issuance of State regulations.

Approved October 29, 2002.

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*LEGISLATIVE HISTORY—s. 2558:*  
*Congressional record, Vol. 148 (2002):*  
*Aug. 1. considered and passed Senate.*  
*Oct 10. considered and passed House.*