

HOUSE No. 2156

The Commonwealth of Massachusetts

In the Year Two Thousand Nine

An Act providing for the establishment and operation of a cancer drug repository program..

Be it enacted by the Senate and House of Representatives in General Court assembled, and by the authority of the same, as follows:

1 SECTION 1. Chapter 111 of the General Laws, as appearing in the 2004 Official
2 Edition, is hereby amended by adding the following section: Section 25J. Cancer drug
3 repository program.

4 Section 1. Definitions. The following words as used in this chapter, unless a different
5 meaning is required by the context or is specifically prescribed, shall have the following
6 meanings:

7 “Board”, the Board of Registration in Pharmacy.

8 “Cancer drug”, a prescription drug that is used to treat any of the following:

9 Cancer or the side effects of cancer.

10 The side effects of a prescription drug used to treat cancer or its side effects.

11 “Commissioner”, the Commissioner of the Department of Public Health.

12 “Department”, the Department of Public Health.

13 “Dispense”, has the meaning given to that term in Section 1 of Chapter 94 C of the
14 General Laws.

15 “Medical facility” means any of the following:

16 An entity licensed by the Commonwealth to provide clinically related health
17 services, including any hospital or clinic, nursing home or long-term care
18 nursing facility, or hospice, as defined in Section 71 of Chapter 111 of the
19 General Laws.

20 A cancer treatment center where radiation therapy and chemotherapy are administered on
21 an ambulatory basis.

22 An office where a physician licensed to practice medicine in the Commonwealth
23 conducts the practice of medicine.

24 “Pharmacy”, any pharmacy licensed by the Commonwealth for transacting retail drug
25 business in accordance with Section 38 of Chapter 112 of the General Laws.

26 “Program”, the Cancer Drug Repository Program established in Section 2.

27 “Supplies”, any supplies used in the administration of a cancer drug.

28 Section 2. Establishment. The Department and the Board of Registration in Pharmacy
29 shall establish a cancer drug repository program for the purpose of authorizing and facilitating
30 the donation of cancer drugs or supplies and their dispensation to residents of the
31 Commonwealth who meet the eligibility standards established in the administrative regulations

32 promulgated by the Department and the Board pursuant to Section 5 of this Act. Under the
33 program, donations may be made on the premises of a medical facility or pharmacy that elects to
34 participate in the program and meets the requirements specified in Section 3 of this Act as well
35 as those promulgated by the Department in regulations.

36 Section 3. Requirements for participation by pharmacies and medical facilities. (a)
37 Participation in the program shall be voluntary.

38 (b) To be eligible for participation in the cancer drug repository program, a pharmacy of
39 medical facility must be licensed and in compliance with all applicable federal and state laws and
40 administrative rules and regulations.

41 (c) A pharmacy or medical facility may fully participate in the cancer drug repository
42 program by accepting, storing, and dispensing donated drugs and supplies, or may limit its
43 participation to only accepting and storing donated drugs and supplies. If a pharmacy or medical
44 facility chooses to limit its participation, the pharmacy or facility shall distribute any donated
45 drugs to a fully participating cancer drug repository in accordance with Section 3.

46 (d) A pharmacy or medical facility may withdraw from participation in the cancer drug
47 repository program at any time upon notification in writing to the commissioner.

48 Section 3. Donation of cancer drugs. Any person or entity, including but not limited to,
49 a cancer drug manufacturer, or medical facility, may donate cancer drugs or supplies to the
50 program or to a participating pharmacy or medical facility. Cancer drugs or supplies may not be
51 donated to a specific cancer patient, and donated drugs or supplies may not be resold by the
52 program or a participating pharmacy or medical facility.

53 Section 4. Acceptance and dispensation of cancer drugs. A cancer drug or supplies
54 needed to administer a cancer drug may be accepted and dispensed under the program if all of
55 the following requirements are met:

56 (a) The cancer drug or supplies to administer a cancer drug is in its original, unopened,
57 sealed and tamper-evident unit-dose packaging or, if packaged in single-unit doses, the single-
58 unit-dose packaging is unopened.

59 (b) The cancer drug bears an expiration date that is later than 6 months after the date that
60 the drug was donated.

61 (c) The cancer drug or supplies needed to administer a cancer drug are not adulterated or
62 misbranded, as determined by a pharmacist who must inspect the drug or supplies before it can
63 be dispensed.

64 The drugs accepted under the program may be distributed to another participating
65 physician's office, pharmacy, hospital or health clinic for dispensing.

66 Nothing in this section shall be construed as prohibiting a pharmacy or medical facility
67 from accepting drugs that are not eligible to be dispensed under the drug repository program for
68 the proper disposal of those drugs.

69 Section 5. Storage, distribution and fees. (a) A pharmacy or medical facility that accepts
70 donated cancer drugs under the program shall comply with all applicable provisions of Federal
71 and State law relating to the storage, distribution and dispensing of cancer drugs.

72 (b) The cancer drugs dispensed through the program shall only be dispensed pursuant to a
73 prescription issued by a prescribing practitioner.

74 (c) A pharmacy or medical facility may charge a handling fee for distributing or
75 dispensing cancer drugs under the program. The fee shall be established in regulations
76 promulgated by the Board.

77 Section 6. Immunity. (a) Unless a pharmaceutical manufacturer exercises bad faith, the
78 manufacturer is not subject to criminal or civil liability for injury, death, or loss to a person or
79 property for matters related to the donation, acceptance, or dispensing of a cancer drug or supply
80 manufactured by the manufacturer that is donated by any person under this Act, including
81 liability for failure to transfer or communicate product or consumer information or the expiration
82 date of the donated cancer drug or supply.

83 (b) Except as provided in Section 6(c), a pharmacy or medical facility participating in the
84 program, a pharmacist dispensing a drug or supply pursuant to the program, a practitioner
85 administering a drug or supply pursuant to the program, or the donor of a cancer drug or supply
86 is immune from civil liability for an act or omission that causes injury to or the death of an
87 individual to whom the cancer drug is dispensed and no disciplinary action shall be taken against
88 a pharmacist or practitioner so long as the drug or supply is donated, accepted, distributed, and
89 dispensed in accordance with the requirements of this section.

90 (c) The immunity or the prohibition of a finding of guilty of unprofessional conduct
91 under Section 6(b) does not extend to the donation, acceptance, distribution, or dispensation of a
92 cancer drug or supply by a person whose act or omission involved reckless, wanton, or
93 intentional misconduct, or malpractice unrelated to the quality of the cancer drug or supply.

94 SECTION 7. In consultation with the Department, the Board shall establish rules and
95 regulations governing the drug repository program that establish all of the following:

96 Eligibility criteria for pharmacies and medical facilities to receive and dispense donated
97 drugs under the program.

98 Standards and procedures for accepting, safely storing, and dispensing donated cancer
99 drugs or supplies needed to administer cancer drugs.

100 Standards and procedures for inspecting donated cancer drugs or supplies needed to
101 administer cancer drugs to determine whether the drugs or supplies are in their original,
102 unopened sealed and tamper-evident unit-dose packaging or, if packaged in single-unit doses, the
103 single-unit-dose packaging is unopened.

104 Standards and procedures for inspecting donated cancer drugs or supplies needed to
105 administer cancer drugs to determine that the drugs or supplies are not adulterated or misbranded
106 and are safe and suitable for dispensing;

107 Eligibility criteria for individuals to receive donated cancer drugs or supplies needed to
108 administer cancer drugs. The standards shall include a method to determine priority of eligible
109 patients under the program.

110 A means, such as an identification card, by which an individual who is eligible to receive
111 donated cancer drugs or supplies needed to administer cancer drugs may demonstrate eligibility
112 to the pharmacy, hospital, or nonprofit clinic dispensing the drugs;

113 A form that an individual receiving a drug from the repository must sign before receiving
114 the drug to confirm that the individual understands the immunity provisions of the program;

115 A formula to determine the handling fee that pharmacies, hospitals, and nonprofit clinics
116 may charge to drug recipients to cover restocking and dispensing costs;

117 A list of cancer drugs and supplies to administer cancer drugs, arranged either by
118 category or by individual drug or supply, that the repository will accept for dispensing.

119 A list of cancer drugs and supplies to administer cancer drugs, arranged either by
120 category or by individual drug or supply, that the repository will not accept for dispensing. The
121 list must include a statement that specifies the reason that the drug or supplies are ineligible for
122 donation.

123 Any other standards and procedures the Board considers appropriate.