

Federal Property Management Regulations

§ 101-42.1102-3

public airports, the Department of Defense or the Federal Aviation Administration, respectively, shall obtain the following signed warning and certification from the donee. State agencies for surplus property shall have the warning and certification typed or stamped on the face of each copy of the distribution document and signed and dated by the authorized representative of the donee organization at the time the property is issued.

Warning and certification:

The donee is aware that the item(s) listed as containing polychlorinated biphenyls (PCBs), a toxic environmental contaminant, require(s) special handling and disposal in accordance with U.S. Environmental Protection Agency regulation (40 CFR part 761) and U.S. Department of Transportation regulations codified in 49 CFR parts 171-180. The donee certifies that this item will be handled and disposed of in accordance with applicable Federal statutes and regulations and applicable State laws.

(d) *Sales requirements.* (1) Surplus PCBs or PCB items normally shall not be sold by GSA or holding agencies. These items are regarded as extremely hazardous and are to be disposed of by the holding agency under the Environmental Protection Agency regulations.

(2) Agencies may request the authority to sell, or that GSA sell, a specific PCB or PCB item. Such requests shall cite the provision in 40 CFR part 761 that authorizes sale and continued use of the specific item. Any such requests shall also include a justification for sale of the item rather than disposal under the EPA regulations.

(3) If PCBs or PCB items are to be sold, the corresponding invitation for bids (IFB), any Standard Form (SF) which lists such items, and any printed matter which advertises the sale of such items shall contain the warning as provided in paragraph (a)(4) of this section.

(e) *Abandonment and destruction.* (1) PCBs and PCB items of personal property not disposed of via utilization, donation, or sale shall be destroyed or otherwise disposed of in accordance with the Environmental Protection Agency regulation (40 CFR part 761) and applicable State laws.

(2) Holding agencies shall contact the nearest office of the EPA for assistance

in complying with the provisions of 40 CFR part 761.

§ 101-42.1102-3 Controlled substances.

(a) *Utilization requirements.* (1) Excess controlled substances are not required to be reported to GSA, but are subject to the utilization screening requirements of § 101-43.311-2. Holding agencies shall make reasonable efforts to obtain utilization of excess controlled substances by offering them to those Federal agencies which certify that they are registered with the Drug Enforcement Administration (DEA), Department of Justice, and are authorized to procure the particular controlled substances requested for transfer. The certification shall include the registration number on the DEA Form 223, Certificate of Registration, issued by DEA.

(2) Holding agencies shall arrange for transfers of controlled substances under §§ 101-43.309-5 and 101-42.207.

(3) All controlled substances that a holding agency determines to be excess shall become surplus after the holding agency has complied with the utilization requirements of paragraph (a)(1) of this section.

(b) *Donation requirements.* Controlled substances shall not be donated.

(c) *Sales requirements.* Surplus controlled substances which are not required to be destroyed as provided in paragraph (d) of this section may be offered for sale by sealed bid under subpart 101-45.3 provided:

(1) The invitation for bids (IFB):

(i) Consists only of surplus controlled substances;

(ii) Requires the normal bid deposit prescribed in § 101-45.304-10;

(iii) Is distributed only to bidders who are registered with the DEA, Department of Justice, to manufacture, distribute, or dispense the controlled substances for which the bid is being submitted; and

(iv) Contains the following special condition of sale:

The bidder shall complete, sign, and return with his/her bid the certificate as contained in this invitation. No award will be made or sale consummated until after this agency has obtained from the Drug Enforcement Administration, Department of Justice, verification that the bidder is registered to manufacture, distribute, or dispense those

§ 101-42.1102-4

41 CFR Ch. 101 (7-1-02 Edition)

controlled substances which are the subject of the award.

(2) The following certification shall be made a part of the IFB (and contract) to be completed and signed by the bidder and returned with the bid:

The bidder certifies that he/she is registered with the Drug Enforcement Administration, Department of Justice, as a manufacturer, distributor, or dispenser of the controlled substances for which a bid is submitted and that the registration number is _____.

Name of bidder (print or type)

Signature of bidder

Address of bidder (print or type)

City, State, Zip code

(3) As a condition precedent to making an award for surplus controlled substances, the following shall be submitted to the Drug Enforcement Administration (DEA), Department of Justice, Washington, DC 20537, Attn: Regulatory Support Section (ODR):

(i) The name and address of the bidder(s) to whom an award is proposed to be made and the bidder(s) registration number(s);

(ii) The name and address of both the holding activity and the selling activity;

(iii) A description of the controlled substances, how those substances are packaged, and the quantity of substances proposed to be sold to the bidder;

(iv) The identification of the IFB by its number, and date on which such bid(s) expire(s); and

(v) A request for advice as to whether the bidder is a registered manufacturer, distributor, or dispenser of controlled substances.

(d) *Destruction of controlled substances.* Controlled substances shall not be abandoned, and destruction of controlled substances must be accomplished in accordance with the terms and conditions applicable to drugs, biologicals, and reagents under §101-42.1102-5(d).

(1) The following shall be destroyed by the holding agency or State agency:

(i) Controlled substances determined surplus at one time and one place with an acquisition cost of less than \$500;

(ii) Controlled substances in a deteriorated condition or otherwise unusable;

(iii) Controlled substances for sale in accordance with §101-42.1102-3(c) but for which no satisfactory or acceptable bids were received.

(2) In addition to the requirements set forth herein, each executive agency and State agency shall comply with the DEA regulations, 21 CFR 1307.21, which provide procedures for disposing of controlled substances, or with equivalent procedures approved by DEA.

(3) Destruction of controlled substances shall be performed by an employee of the holding agency or State agency in the presence of two additional employees of the agency as witnesses to that destruction unless the special agent in charge (SAC) of the DEA Divisional Office directs otherwise.

§ 101-42.1102-4 Nuclear Regulatory Commission-controlled materials.

(a) *General.* The Nuclear Regulatory Commission (NRC) has exclusive control over licensing, use, transfer, and disposition of NRC-controlled materials.

(b) *Transfer of NRC-controlled materials.* NRC-controlled materials shall not be reported to GSA as excess personal property, nor shall they be made available for excess and surplus screening as nonreportable property. Transfer and disposition of such materials do not require GSA approval and shall be accomplished only under the applicable regulations of the NRC (see 10 CFR parts 30 through 35, 40, and 70).

(c) *Information and inquiries.* All inquiries for further information or specific instructions regarding the licensing, use, transfer, or disposition of NRC-controlled materials shall be directed to the U.S. Nuclear Regulatory Commission, Washington, DC 20555.

§ 101-42.1102-5 Drugs, biologicals, and reagents other than controlled substances.

In addition to the requirements of subparts 101-42.2 through 101-42.4, drugs, biologicals, and reagents which are fit for human use shall be reported as provided in this §101-42.1102-5. Drugs, biologicals, and reagents that