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