

Misc expired medical supplies (368370527510021) One lot consisting of 2 full pallets of expired medical items. Too many individual items to list. Catheters, Tray Lumbar Puncture, Kit Vascular Cath, Peel-Away Introducer Set; Tube, NG; Set, Suprapubic Catheter Dispos; Endopath Surgical Thoracic Torcar sleeve w/rounded tip; MILD device kit for LSS treatment lumbar decompressive procedures; Glove Surg Size 9; Glove Surg Size 8; Arterial Safety valve Retroguard; Huber needles safe set; Pneumoneedle Insufflation w/luer lock; Central Venous Catherization 16ga; Exchange guide wires; \*\*All items are expired\*\* Sale intended for training/veterinary use only. Winning bidder required to complete and submit the attached "71QSCI16019013 Med Devices SOI.pdf" prior to removal, e-mail to mark.maxwell@gsa.gov is the preferred method for submission of that document.

Purchasers of all medical equipment listed in the Invitation for Bid (IFB) shall certify and assure in writing that such item will be used or resold only under the conditions specified below:

Medical device items are subject to the laws and regulations administered by the Food and Drug Administration (FDA). Provisions of the governing statute, the Federal Food, Drug and Cosmetic Act appear in 21 U.S.C. 331, ET. Seq. In summary, the Act prohibits the movement in interstate commerce of medical devices that are misbranded or adulterated. The Act authorizes FDA to initiate criminal enforcement proceedings against companies and/or individuals responsible for violations of its provisions. Moreover, the Act authorizes FDA to initiate civil proceedings to seize, or enjoin the distribution of such items.

It shall, also, be the responsibility of all purchasers to comply with local, state, or other applicable laws.

The following certificate, to be a separate attachment to the Invitation for Bid, is required by FDA to purchase the medical device items identified in the Invitation.

I certify that I am a licensed practitioner and/or other person regularly and lawfully engaged in the manufacture and/or refurbishing of the medical device item identified in the IFB. I, also, certify that prior to sale or use of such a device, I will take assurance that such a device is not adulterated or misbranded within the meaning of those terms in the Federal Food Drug and Cosmetic Act (21 U.S.C., et Seq.).

\_\_\_\_\_  
**Signature**

\_\_\_\_\_  
**Date**

Recognizing that Federal law places stringent restrictions on adulterated or misbranded medical devices (21 U.S.C. 331, et. Seq.), I certify that I either will sell or otherwise proffer the medical device item identified in the IFB to persons described in the above, or will not use this item(s) for their original or usual intended use, for any other medical use.

\_\_\_\_\_  
**Signature**

\_\_\_\_\_  
**Date**

False or misleading statements may result in a fine of not more than \$10,000 or imprisonment for not more than five (5) years, or both (18 U.S.C. 1001).