CHANGES TO AATB STANDARDS FOR TISSUE BANKING
14th Edition

SECTION D
AUTHORIZATION, INFORMED CONSENT, DONOR SCREENING, AND TISSUE RECOVERY, COLLECTION, AND ACQUISITION

Current (14th Edition)
D5.000 RECOVERY, COLLECTION, AND ACQUISITION

D5.100 Reagents, Supplies, Materials, and Equipment
All *critical* supplies, reagents, materials, and equipment approved for use for *recovery, collection,* or *acquisition shall* be identified and specifications (e.g., *sterile* where applicable) documented. A *record shall* be made of all reagents, supplies, and materials following receipt including, as applicable, the type, quantity, manufacturer, *lot* number, date of receipt, and expiration date or manufacturing date (as applicable). Inspection *shall* be documented, including identification of the staff performing the inspection. The *tissue bank shall* maintain *records* of all supplies, reagents, materials, and equipment from receipt through period of time used. All reagents, supplies, materials and equipment *shall* be used and stored in accordance with manufacturers' instructions, unless qualified/*validated* for intended use or storage.

All non-disposable surgical instruments and parts of mechanical/ electrical equipment which come in contact with *tissue shall* be properly cleaned, *decontaminated*, and *sterilized* prior to use for *recovery, collection,* or *acquisition* according to written *procedures* prepared to prevent contamination or *cross-contamination*. *Records shall* be maintained that document sterilization steps. All reagents, supplies, and materials *shall* be used and stored in accordance with manufacturers' instructions unless qualified/validated for intended use or storage.

With Amendments
D5.000 RECOVERY, COLLECTION, AND ACQUISITION

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As Amended

D5.000 RECOVERY, COLLECTION, AND ACQUISITION

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