premarket notification procedures in subpart E of part 807 of this chapter subject to §864.9.


§864.9875 Transfer set.

(a) Identification. A transfer set is a device intended for medical purposes that consists of a piece of tubing with suitable adaptors used to transfer blood or plasma from one container to another.

(b) Classification. Class II (performance standards).

[45 FR 60651, Sept. 12, 1980]

Subpart K—Products Used In Establishments That Manufacture Human Cells, Tissues, and Cellular and Tissue-Based Products (HCT/Ps)

§864.9900 Cord blood processing system and storage container.

(a) Identification. A cord blood processing system and storage container is a device intended for use in the processing and the storage of cord blood. This device is a functionally closed processing system that includes containers, other soft goods, and a centrifugation system for cord blood concentration, and a final container for the cryopreservation and the storage of a cord blood product.

(b) Classification. Class II (special controls). The special control for this device is FDA’s guidance document entitled “Class II Special Controls Guidance Document: Cord Blood Processing System and Storage Container.” For the availability of this guidance document, see §864.1(d).

[72 FR 4638, Feb. 1, 2007]

PART 866—IMMUNOLOGY AND MICROBIOLOGY DEVICES

Subpart A—General Provisions

Sec. 866.1 Scope.
866.3 Effective dates of requirement for premarket approval.
866.9 Limitations of exemptions from section 510(k) of the Federal Food, Drug, and Cosmetic Act (the act).

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Subpart B—Diagnostic Devices

866.1620 Antimicrobial susceptibility test disc.
866.1640 Antimicrobial susceptibility test powder.
866.1645 Fully automated short-term incubation cycle antimicrobial susceptibility system.
866.1700 Culture medium for antimicrobial susceptibility tests.

Subpart C—Microbiology Devices

866.2050 Staphylococcal typing bacteriophage.
866.2120 Anaerobic chamber.
866.2160 Coagulase plasma.
866.2170 Automated colony counter.
866.2180 Manual colony counter.
866.2300 Multipurpose culture medium.
866.2320 Differential culture medium.
866.2330 Enriched culture medium.
866.2350 Microbiological assay culture medium.
866.2360 Selective culture medium.
866.2390 Transport culture medium.
866.2410 Culture medium for pathogenic Neisseria spp.
866.2420 Oxidase screening test for gonorrhea.
866.2440 Automated medium dispensing and stacking device.
866.2450 Supplement for culture media.
866.2480 Quality control kit for culture media.
866.2500 Oxidase screening test for gonorrhea.
866.2540 Microbiological incubator.
866.2560 Microbial growth monitor.
866.2580 Gas-generating device.
866.2600 Wood’s fluorescent lamp.
866.2660 Microorganism differentiation and identification device.
866.2850 Automated zone reader.
866.2900 Microbiological specimen collection and transport device.

Subpart D—Serological Reagents

866.3010 Acinetobacter calcoaceticus serological reagents.
866.3020 Adenovirus serological reagents.
866.3035 Arizona spp. serological reagents.
866.3040 Aspergillus spp. serological reagents.
866.3050 Beta-glucan serological assays.
866.3060 Blastomyces dermatitidis serological reagents.
866.3065 Bordetella spp. serological reagents.
866.3085 Brucella spp. serological reagents.
866.3110 Campylobacter fetus serological reagents.
866.3120 Chlamydia serological reagents.
866.3125 Citrobacter spp. serological reagents.
866.3135 Coccidioides immitis serological reagents.
866.3140 Corynebacterium spp. serological reagents.