

“GUIDANCE FOR INDUSTRY: RECOMMENDATIONS FOR OBTAINING A LABELING CLAIM FOR COMMUNICABLE DISEASE DONOR SCREENING TESTS USING CADAVERIC BLOOD SPECIMENS FROM DONORS OF HUMAN CELLS, TISSUES, AND CELLULAR AND TISSUE-BASED PRODUCTS (HCT/PS);” Availability

DOCKET NO. 2005D-0056

REFERENCES

- TAB 1 Human Tissue Intended for Transplantation; Final Rule, July 29, 1997 (62 FR 40429), (<http://www.fda.gov/cber/genadmin/frtissue.txt> or <http://www.fda.gov/cber/tissue/docs.htm>).
- TAB 2 Guidance for Industry: Screening and Testing of Donors of Human Tissue Intended for Transplantation; Notice of Availability, July 29, 1997 (62 FR 40536), (<http://www.fda.gov/cber/gdlns/tissue2.txt> or <http://www.fda.gov/cber/tissue/docs.htm>).
- TAB 3 Guidance for Industry: Availability of Licensed Donor Screening Tests Labeled for Use with Cadaveric Blood Specimens, June 2000, (<http://www.fda.gov/cber/gdlns/cadbld.htm> or <http://www.fda.gov/cber/tissue/docs.htm>).
- TAB 4 Eligibility Determination for Donors of Human Cells, Tissues, and Cellular and Tissue-Based Products; Final Rule, May 25, 2004 (69 FR 29786), (<http://www.fda.gov/cber/tissue/docs.htm>).
- TAB 5 Tissue related documents from CBER Web Site

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